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SUGARBEET RESEARCH

1999 REPORT



FOREWORD

SUGARBEET RESEARCH is an annual compilation of progress reports concerning research by U. S. Department of Agriculture, Agricultural Research Service investigators and other cooperators who are engaged in sugarbeet research. The report was assembled and produced at the expense of the Beet Sugar Development Foundation, and is for the sole use of its members and the cooperators. Much of the data has not been sufficiently confirmed to justify general release and interpretations may be modified with additional experimentation. This report is not intended for publication and should not be used for cited reference nor quoted in publicity or advertising. Reproduction of any portion of the material contained herein will not be permitted without the specific consent of the contributor and the Beet Sugar Development Foundation.

The report presents results of investigations strengthened by contributions received under Cooperative Agreement between the USDA Agricultural Service and the Beet Sugar Development Foundation, along with the California Beet Growers Association, the Western Joint Research Committee, the Sugarbeet and Education Board of Minnesota and North Dakota, and Texas A & M University.

Trade names occur in this report solely to provide specific information and do not signify endorsement by the U. S. Department of Agriculture, Texas A & M University, the Beet Sugar Development Foundation or any of the cooperating organizations.

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SUGARBEET RESEARCH

1999 REPORT

Section A

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ABSTRACTS OF PAPERS PUBLISHED OR APPROVED FOR PUBLICATION, 1999

LEWELLEN, R.T. <u>Registration of Multiple Disease Resistant C69 Sugarbeet Germplasm</u>. Crop Sci. 40 (in press).

Sugarbeet (Beta vulgaris L.) germplasm line C69 (PI599341) was developed by the USDA-ARS in cooperation with the Beet Sugar Development Foundation and the California Beet Growers Association. This line was released in 1997. C69 is a vigorous, multigerm, self-sterile line with tolerance to virus yellows (VY) and segregates for resistance to rhizomania conditioned by Rz. Tolerance to VY is to beet western yellows, beet chlorosis, and beet yellows viruses, but resistance to the luteoviruses, beet western yellows and beet chlorosis, is stronger. The VY tolerance was derived from nearly the entire germplasm base of the long term USDA-ARS VY resistance breeding program at Salinas, CA. The Rz allele was from lines C78, C79, C80, and C82 that were developed by backcrossing Rz into C46, C37, C54, and C31, respectively. Plants selected for resistance to rhizomania caused by beet necrotic yellow vein virus within lines C78, C79, C80, and C82 were bulked and used as the pollinator in a composite cross made in a field seed plot to rhizomania susceptible stecklings combined from lines C31/6,C39, C46/2, C47, C49, C54, C91, C92, Y48, Y56, and Y57. Plants selected for resistance to rhizomania from this composite cross were bulked and used to pollinate a composite of stecklings with green hypocotyls from breeding lines C31/6, C31-43, C31-89, C39, C49, and C91. Both red hypocotyl color and resistance to rhizomania were used as markers to identify F₁ plants from this second composite cross. The inter se increase of 71 F₁ plants produced a broadly-based line called Y569. Y569 is expected to be predominantly composed of the germplasm of C31 with smaller amounts of C37, C46, C39, C64, and other sources.

Y569 was planted under moderately severe rhizomania conditions at Salinas. The plants in the selection plot were inoculated with sugarbeet *Erwinia* [*E. carotovora* (Jones) Bergey subsp. *betavasculorum*]. Powdery mildew caused by *Erysiphe polygoni* DC and Cercospora leaf spot caused by *C. beticola* Sacc. were not controlled and were moderate on susceptible plants. A high incidence of natural infection with beet western yellows virus occurred. Phytophthora tip rot caused by *P. dreschleri* Tucker was prevalent and differentially damaged breeding lines in this planting. In late November, individual plants were selected based upon root yield, beet conformation, and resistance to rhizomania, root rots, Cercospora leaf spot, and powdery mildew. Roots visually selected in the field were reselected for sucrose concentration. Following vernalization, 39 (about 2% of initial population) mother roots were increased in mass to produce breeding line Y769. Line Y769 was reselected for resistance to rhizomania to produce germplasm C69.

Preliminary tests show that C69 has relatively high sucrose concentration, good root and agronomic traits, large canopy, and combined disease resistance. Segregation for reaction to rhizomania occurs. This line has high resistance to *Erwinia* and moderate resistance to VY and powdery mildew. In tests under VY inoculated conditions, C69 had higher sugar yield than any

breeding line or commercial hybrid in the trials. It is a moderately nonbolting type. Line C69 is moderately susceptible to curly top virus. It has higher sucrose concentration than C78 or C80. Line C69 should be useful as a broadbased source for continued population improvement and from which parental lines with combined resistance could be extracted.

LEWELLEN, R.T. <u>Registration of Powdery Mildew Resistant Sugarbeet Germplasms CP01</u> and CP02. Crop Sci. 40 (in press).

Sugarbeet (Beta vulgaris L.) germplasm lines CP01 (PI610490) and CP02 (PI610491) were developed by the USDA-ARS in cooperation with the Beet Sugar Development Foundation and the California Beet Growers Association. These lines were released in 1999. CP01 and CP02 are self-sterile, multigerm, germplasm lines that segregate for resistance to powdery mildew caused by Ervsiphe polygoni DC (syn. E.betae Weltzien). CP01 and CP02 have identical developmental histories except for the source of resistance to powdery mildew. Resistance within CP01 was from WB97 (PI546394) and within CP02 was from WB242 (PI546413). High resistance to powdery mildew within these B. vulgaris L. subsp. maritima (L.) Arcang. accessions was identified separately by McFarlane and Whitney at Salinas, CA. Seed of WB97 was sent to Salinas from the Japan Sugarbeet Improvement Foundation in 1968. Passport information indicated that it was sent to Japan from Wageningen, the Netherlands as WB46 (B.patula Ait.) in 1963. The site of its original collection is not known. If WB46 is B.patula, then it would have been collected from Ilheu dos Embarcaderos near Madeira. Seed of WB242 was obtained from IRS, Bergen op Zoom, the Netherlands, in 1974 as a B. vulgaris subsp. maritima line originally collected in the Loire River Estuary in France. WB242 also is known to have low sugarbeet cyst nematode (SBCN) Heterodera schachtii Schmidt counts and may be the same or similar to wild beet lines known as Le Pouliguen Group 2 and to PI198758 and PI198759. When grown at Salinas, WB242 is variable for plant type, root and stem pigmentation, bolting habit, and root type. Most plants of both WB97 and WB242 have red hypocotyls and stems and are annual. Both are susceptible to rhizomania caused by beet necrotic yellow vein virus.

In order to enhance sugarbeet with the high resistance to powdery mildew found in WB97 and WB242 and to study the inheritance of powdery mildew resistance, powdery mildew resistance was backcrossed into sugarbeet breeding line C37 that has resistance to curly top, virus yellows, *Erwinia* sp. and bolting. C37 is highly susceptible to powdery mildew, completely self-sterile under Salinas greenhouse conditions, and has green hypocotyls. These traits facilitated making and recognizing the F₁ hybrids in each generation. Resistance from WB97 and WB242 was transferred in separate but parallel sets of crosses. Usually C37 was used as the female parent so CP01 and CP02 have sugarbeet cytoplasm. CP01 and CP02 initially were released as the BC₄F₂ generation. BC₄F₁ testcross families of these lines were evaluated in the field in 1997 and segregated for reaction to powdery mildew. Unselected stecklings of these BC₄F₁ testscrosses were increased in mass to produce lines P813 and P814 that were released as CP01 and CP02, respectively. Genetic studies in 1997 and 1999 indicated that resistance to powdery mildew is inherited in the manner of a single dominant allele in each of these wild beet sources. This resistance has tentatively been assigned the *Pm* gene symbol. Allelism between the WB97 and WB242 resistances has not been determined.

CP01 and CP02 are susceptible to rhizomania. Likewise, they should be similar to the C37 recurrent parent for other traits. Some of the BC₄F₁ testcrosses segregated for annualism so this trait remains in these lines. No attempt has been made to determine if any variability for SBCN resistance remains from WB242. CP01 and CP02 should be useful as enhanced sources of resistance to powdery mildew originally found in *B.vulgaris* subsp. *maritima* and for genetic research. Preliminary tests have tentatively identified molecular markers specific for powdery mildew resistance from WB242.

OBERMEIER, C., J.L. SEARS, G.C. WISLER, H.Y. LIU, K.O. SCHLUETER, E.J. RYDER, J.E. DUFFUS, and S.T. KOIKE. 1999. <u>Characterization of a new tomato bushy stunt-related tombusvirus associated with lettuce dieback disease in California</u>. Phytopathology 85:S57.

A disease causing dieback of Romaine lettuce has been found increasingly in California. Affected lettuce plants exhibit severe stunting, chlorosis and necrosis of older leaves. Plants infected early in their development may die. An isometric virus has been isolated consistently from roots and leaves of symptomatic lettuce plants. Particles are 30 nm in diameter. Double-stranded RNA profiles are identical to those of TBSV isolates. Cloning of the 3í-terminus of the viral genomic RNA revealed 84% to 88% nucleotide sequence identity with several TBSV strains. RT-PCR has been successfully applied for detection of the virus in lettuce leaves. Field trials revealed no resistance in Romaine, but did show resistance in several leaf and crisphead lettuce varieties. Although inoculation under greenhouse conditions has not yet reproduced the dieback disease in lettuce, the consistent isolation of this TBSV-related virus from field-grown symptomatic lettuce suggests that it may be the cause of the disease.

WEILAND, J.J. and R.T. LEWELLEN. Generation of Molecular Genetic Markers Associated with Resistance to Powdery Mildew (*Erysiphe polygoni* DC) in Sugarbeet (*Beta vulgaris* L.). 9th Int'l Congress, July 1999. International Soc. Plant-Microbe Interactions. p. 215

Powdery mildew caused by *Erysiphe polygoni* DC can be devastating to sugarbeet production particularly in warm, dry climates. Although resistance to certain races of *E.polygoni* exists in sugarbeet, powdery mildew disease is typically controlled through the use of fungicides. The identification of broad resistance to sugarbeet powdery mildew in the wild beet *B. vulgaris* spp. *maritima* was followed by the incorporation of this resistance into sugarbeet by recurrent backcrossing and progeny testing. Germplasm accession C37 was used as the susceptible, recurrent parent and P604 is the F₂BC₃ population at the intermediate stage of the introgression. Three DNA pools each were produced for C37 and P604; each pool was comprised of the DNA from 7 individual plants. A diprimer adaptation of the RAPD analysis was applied to the DNA pools, where one of the primers was composed of a sequence homologous to that encoding a core sequence found in many plant disease resistance genes. Amplified products were identified that were associated with all three DNA pools derived from P604 plants, but with none of the three DNA pools derived from C37. The possibility that some of the amplified products contain sequences of the gene conferring resistance to sugarbeet powdery mildew is discussed.

WINTERMANTEL, W.M. and J.L. SEARS. 1999. <u>Examination of viral interactions in relation to disease severity and resistance in the virus yellows complex of sugarbeet</u>. Phytopathology 89:S85.

Virus yellows is a disease complex composed of different genera of plant viruses. Beet yellows closterovirus (BYV), beet western yellows luteovirus (BWYV), and occasionally, beet mosaic potyvirus (BtMV), are the main components. BtMV alone may not contribute to economically significantly disease loss. All of these viruses are transmitted by aphids, and all are usually present at some level in infected fields. Although beet-free periods are useful in managing virus yellows, the increased range and population of the black bean aphid has made this disease more difficult to control in recent years. In this study, sugarbeet varieties exhibiting differential levels of resistance to the yellows complex viruses were inoculated with every possible combination of one, two or all three viruses. Interviral effects were identified and correlated using quantitative molecular techniques. Correlation of stunting and symptom severity with different virus combinations indicate that disease is more severe when all three viruses are present than when plants are infected by one or any combination of two viruses.

WINTERMANTEL, W.M. AND M. ZAITLIN. 2000. <u>Transgene translatability increases</u> <u>effectiveness of replicase-mediated resistance to Cucumber mosaic virus</u>. J. Gen. Virol. 81, 587-595.

Transgenic tobacco plants expressing an altered form of the 2a replicase gene from the Fny strain of cucumber mosaic virus (CMV) exhibit suppressed virus replication and restricted viral movement when inoculated mechanically or by aphid vectors. Additional transformants have been generated which contain replicase gene constructs designed to determine the role(s) of transgene mRNA and/or protein in resistance. Resistance to systemic disease caused by CMV, as well as delayed infection, was observed in several lines of transgenic plants which were capable of expressing either full length or truncated replicase proteins. In contrast, among plants which contained nontranslatable transgene constructs, only one of sixty-one lines examined exhibited delays or resistance. Once infected, plants never recovered, regardless of transgene translatability. Transgenic plants exhibiting a range of resistance levels were examined for transgene copy number, mRNA and protein levels. Although ribonuclease protection assays demonstrated that transgene mRNA levels were very low, resistant lines had consistently more steady-state transgene mRNA than susceptible lines. Furthermore, chlorotic or necrotic local lesions developed on the inoculated leaves of transgenic lines containing translatable transgenes, but not on inoculated leaves of lines containing nontranslatable transgenes. These results demonstrate that translatability of the transgene and possibly expression of the transgene protein itself facilitates replicase-mediated resistance to CMV in tobacco.

WISLER,G.C., R. T. LEWELLEN, W. M. WINTERMANTEL, H-.Y. LIU, and J. S. SEARS. 1999. <u>Differences Among Sugarbeet Cultivars with Varying Levels of Rhizomania Resistance To Single And Mixed Infections with BNYVV and BSBMV</u>. Proc. Fourth Symposium of the International Working Group on Plant Viruses with Fungal Vectors. 135-138.

Eight sugarbeet cultivars, that range in reaction to rhizomania from uniformly susceptible to highly resistant, were compared for levels of beet necrotic yellow vein benyvirus, as measured by TAS-ELISA in field studies in Salinas, California. Differences in absorbance (A_{405 nm}) values measured among the cultivars closely correlated with the dosage and frequency of the Rz allele that conditions resistance to BNYVV. Absorbance values were significantly positively correlated with rhizomania disease index scores and negatively correlated with individual root weight, plot root weight, and sugar yield. The same eight cultivars were compared in greenhouse pot cultures for their reactions to beet soil-borne mosaic benyvirus. All cultivars were highly susceptible to BSBMV, with absorbance readings ranging from 8 to 12 times the healthy root mean. When mixed infections of BNYVV and BSBMV were compared to single infections in a susceptible and resistant sugarbeet line, the reactions, as measured by root symptoms and individual beet weight were significantly more severe than each virus alone. This was true regardless of whether the seedlings were initially infested with either BNYVV or BSBMV. Thus, resistance to BNYVV does not confer resistance to BSBMV, nor does BSBMV infection moderate the effects of BNYVV.

WISLER, G.C., J. L. SEARS, H.-Y. LIU, C. OBERMEIER, and J. E. DUFFUS. 1999. <u>A new disease of greenhouse-grown tomatoes caused by tomato bushy stunt virus (TBSV)</u>. Phytopathology 89:S85.

A previously undescribed disease of hydroponic, greenhouse-grown tomatoes was detected in the Central United States. Symptoms include stunting of affected plants, leaf necrosis, fruit and flower drop, and truss necrosis. Although fruit appears to be normal, the stem end shows a ring of necrosis after the calyx is removed, and the internal part of the fruit shows necrosis that is primarily restricted to the vascular tissues. TBSV has been consistently isolated from symptomatic foliage, trusses and fruit. No fungal or bacterial organism has been isolated from symptomatic tissues. Virus particles measure 30 nm in diameter. The dsRNA profile is identical to those of known TBSV isolates. Koch's postulates were completed by pouring inoculum, increased in *Nicotiana benthamiana* from single local lesions, into 10 cm pots with tomato 'Trust' seedlings. Foliage and truss necrosis was produced by this method, and TBSV was re-isolated from affected tissues. Based on the unique fruit symptoms observed, this may be a different isolate or strain of TBSV than previously identified in tomato.

YU, M.H. Root-knot nematodes in California and the development of resistant sugarbeet varieties. Proc. Agric. Am. Soc. Sugar Beet Technol. p. 167-173. 1999.

The status of root-knot nematode distribution in California sugarbeet fields was investigated. Samples of the galled plants and infested soil were collected from various major growing areas. To identify the specificity of *Meloidogyne* spp., nematodes were initially recovered with the use of susceptible hosts. Matured females and egg masses were extracted from infected plants and inoculated to individual tomato seedlings that were growing in cone-tainers; for nematodes from infested soil, seedlings were germinated directly in pots containing the field soil to induce galling. Isolates recovered from these procedures were increased to build populations. They were then inoculated to groups of test plants for the 'differential host assay'. The results

indicated that the four most common species of root-knot nematode, i.e., *M. incognita*, *M. javanica*, *M. arenaria*, and *M. hapla*, were currently existent in California sugarbeet growing areas, occurring in eleven or more counties. Genetic sources of resistance to root-knot nematode is now available. Due to its multi-species resistance capability, sugarbeet production may be protected from serious root-knot nematode damages when the resistance is eventually incorporated into a commercial variety.

YU, M.H., W. HEIJBROEK and L.M. PAKISH. The sea beet source of resistance to multiple species of root-knot nematode. Euphytica 108: 151-155. 1999.

Development of commercially available host-plant resistance to *Meloidogyne* spp. is essential to sugarbeet (*Beta vulgaris* L.) root-knot nematode resistance breeding. Reactions of seedlings from resistant crosses and hybrid derivatives were evaluated against juvenile (J2) inoculations in the greenhouse. The noncultivated sea beet [B. vulgaris ssp. maritima (L.) Arcang] source of resistance is effective against the four economically important root-knot nematodes, i.e., M. incognita Races 1, 2, and 4 (Race 3 not tested), M. javanica, M. arenaria Races 1 and 2, and M. hapla. In monoxenic culture, M. arenaria inoculations resulted in the most galling, and M. hapla, the least. Species combinations induced higher rates of infection. Different races of the same Meloidogyne species caused similar galling. Preliminary inoculation studies indicated that resistance was also effective to M. chitwoodi and M. fallax. The trait of resistance to multiple Meloidogyne species may be valuable in developing sugarbeet, and possibly transgenic lines of other crops, resistant to root-knot nematode.

Project 281

Evaluation of BNYVV and BSBMV Concentrations and Effects of Rhizomania Resistant and Susceptible Sugarbeet Varieties

G. C. Wisler, R. T. Lewellen, W. M. Wintermantel, H.-Y. Liu, and J. E. Duffus

Rhizomania continues to be an important disease problem for the sugarbeet industry. Our research program regarding the detection and differentiation of Beet necrotic vellow vein virus (BNYVV), the cause of Rhizomania, and related soil-borne viruses belonging to the same genus Benyviridae, (formerly the Furovirus group) has made significant contributions over the past several years. A member of this family of viruses, termed Beet soil-borne mosaic virus (BSBMV), has received an increasing amount of attention due to the fact that it is serologically related to BNYVV. This has caused problems in certain serological tests due to low levels of cross-reactivity. Initial research on this particular project began in 1993 where five BNYVV isolates and eight BSBMV isolates from the United States were compared using serological. molecular and biological tests. It was concluded from these tests that all BNYVV isolates in the United States are virtually identical to one another. In addition, all BSBMV isolates, although serologically identical to one another, differ in plant host reactions and molecular properties. Another important conclusion drawn from this research was that BNYVV was shown to be clearly distinct from BSBMV, and BSBMV was determined to be a distinct benyvirus. These results have been repeatedly confirmed by additional tests in our lab for the past six years and more recently by others.

There are several highly sensitive diagnostic tests on the market today due to the massive screening programs developed for HIV. These tests are based on amplification of portions of the viral genome [by reverse transcriptase-polymerase chain reaction (RT-PCR)] followed by light activated molecular labels that are measured by sensitive equipment. However, these systems are not generally applicable to agricultural research due to the extreme expense of this technology. We have evaluated both amplification of the BNYVV genome by RT-PCR and subsequent detection with labeled molecular probes. We have evaluated several different variations of these methods and compared them to ELISA for BNYVV diagnosis. We have made significant improvements in the ELISA technique based on (1) production of antisera to the cloned coat protein of BNYVV, which provides a long-term supply of identical protein for future immunizations, and (2) using it in a triple-antibody sandwich ELISA in combination with a monoclonal antibody. This modified ELISA significantly improves the test by eliminating the cross-reaction, which was seen between BNYVV and BSBMV, and increases the sensitivity by adjusted concentrations of reagents. This test is now being marketed by Agdia, Inc. (Elkhart, Indiana). Thus, for the purpose of sensitive and specific identification of Rhizomania from soil samples, the TAS-ELISA, which is available from Agdia, is the best choice at this time.

Our studies on the BSBMV isolates of sugarbeet in the United States has suggested that this virus is responsible for some significant yield losses in areas that are infested. Because we have never detected BSBMV in California, we have to study this virus in other states where it is prevalent, or in greenhouses at the USDA in Salinas. BSBMV has been found in several locations in Colorado, Nebraska and Minnesota. In last year's Blue Book we reported results from 27 fields in Colorado and Nebraska where a significant decline in yield has been experienced by growers for the past few years. In that study, no fungal or bacterial pathogen was

found in any sample, and soil analyses indicated normal levels of nutrients, pH, etc. However. 24 of 27 fields were found to be infested with either BSBMV. Beet soil-borne virus (BSBV) another benyvirus member), or both. Only one field was found to be infested with Rhizomania. Although we do not know the exact cause of this yield decline, the association of low yields with these viruses which we know (from greenhouse studies) to be the cause of reduced growth of beets, suggests that we should pursue this line of research. This year we performed a small variety trial in Nebraska where half the plot was fumigated. Although the field had very low levels of BNYVV and BSBMV, a trend was observed where the fumigated section of the field had increased yields and sugar (see Western Sugar annual report). Studies in the past year have focused on determining if the series of Rhizomania resistant varieties used in the previous year's report show any resistance to BSBMV. We did not expect this to be the case, since BNYVV and BSBMV are distinct viruses and resistance would not normally be conferred to more than one virus unless they were considered to be strains or isolates of one another. The varieties used in this test are listed in Table 1. Results from last year's study showed: (1) differences in absorbance values for BNYVV measured by TAS-ELISA among the eight cultivars were closely correlated to the dosage and frequency of the Rz allele that conditions resistance to BNYVV. (2) the diploid Rzrz hybrid Beta4776R had a significantly lower value than the similar triploid Rzrzrz hybrid Beta4038R, and (3) cultivars that segregated Rzrz:rzrz (i.e., SS-781R and 6921H50) had higher absorbance values than the uniformly resistant Rzrz hybrids Beta4776R and HM7072. We also found that the virus titers (concentrations) in infected beets declined throughout the season. Therefore, sampling early in the season gives the best estimate of the disease incidence in the field.

Table 1. Sugar beet hybrids evaluated in rhizomania experiments; Salinas, California, 1997 growing season

Identification	Source	Description	Genotype
USH11	USDA-ARS	diploid susceptible	rzrz
KWS6770	Betaseed	triploid susceptible	rzrzrz
Beta4776R	Betaseed	diploid resistant	Rzrz
SS-781R	Spreckels	diploid segregating	Rzrz:rzrz
Rival	Holly	diploid resistant	Rzrz
HM7072	Novartis	diploid resistant	Rzrz
Beta4038R	Betaseed	triploid resistant	Rzrzrz
6921H50	USDA-ARS	diploid segregating	B. maritima hybrid

The same eight varieties that were used in the Rhizomania field study were also used in greenhouse studies to evaluate their possible resistance to BSBMV. Soil was obtained from fields that had been previously tested and were infested with BSBMV only. All eight varieties showed high BSBMV readings in DAS-ELISA in BSBMV soils (Fig. 1). Thus, it appears that resistance to BNYVV does not confer resistance to BSBMV.

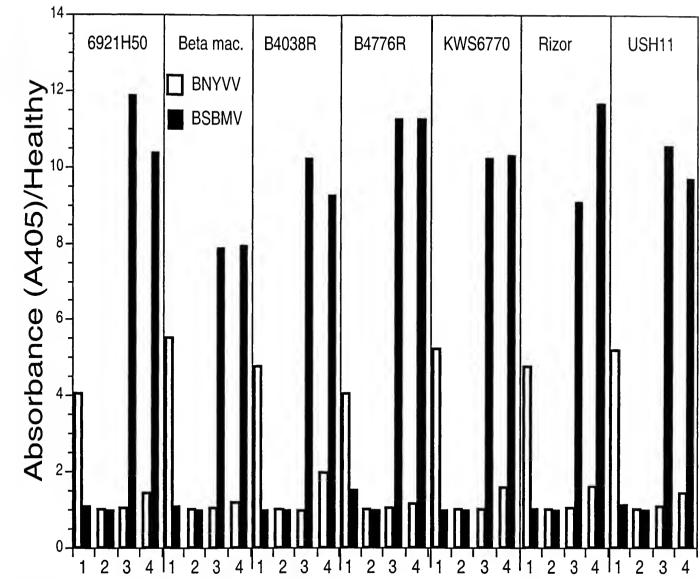


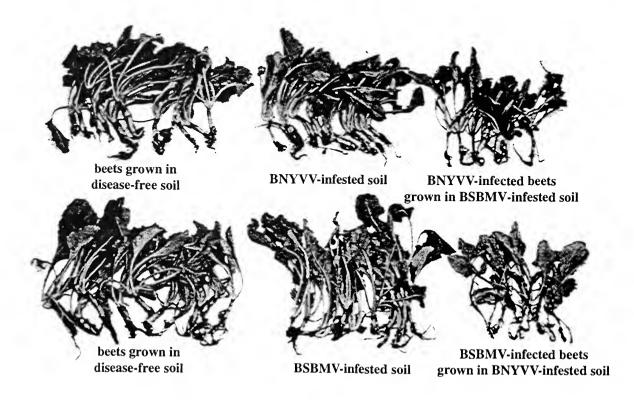
Fig. 1.

Cultivars USH-11 and B4776R were planted in greenhouse studies to determine the effect of mixed infections of BNYVV and BSBMV in sugar beets. Seed was first planted into (i) healthy soil, or soil infested with either (ii) BNYVV, or (iii) BSBMV. After six weeks they were tested for each virus, then transplanted into the respective soil; healthy, BNYVV- or BSBMV-infested. After an additional six weeks, plants were tested again for BNYVV and for BSBMV evaluated for symptoms and roots were weighed (Fig. 2). Only those beets that were infected with BNYVV showed blackened roots. Otherwise, the beets infected with BSBMV only were reduced in weight. Other than general yellowing, no leaf symptoms were observed for either virus. Depending on the variety used, the effect of mixed infections ranged from a 30% fresh weight reduction to an 81% reduction (Table 2). This study shows that under conditions of mixed infections in the soil, neither virus appears to moderate the infection of the other.

Table 2. Results from Cross-inoculation Experiments; Salinas, California, 1999.

Table 2. Results Heart Cross					
		Soil	Mean fresh wt.	BNYVV	BSBMV
Variety	Seedlings	Transplanted	(10 beets)	TAS-ELISA	TAS-ELISA
	O	Into		(OD/Healthy)	(OD/Healthy)
B4776R °	healthy	healthy	101.3g ^a	1.00 ^b	0.976
B4776R	BSBMV	healthy	76.1	1.03	0.964
USH-11°	healthy	healthy	71.22	1.66	1.000
USH-11	BSBMV	healthy	66.7	0.99	0.960
B4776R	BNYVV	healthy	63.6	4.18	1.028
B4776R	healthy	BSBMV	38.97	1.14	3.788
B4776R	healthy	BNYVV	38.35	2.65	0.996
B4776R	BNYVV	BSBMV	29.14	2.87	4.088
B4776R	BSBMV	BNYVV	25.84	3.03	1.108
USH-11	healthy	BNYVV	24.74	4.75	0.968
USH-11	healthy	BSBMV	23.84	1.16	4.064
USH-11	BSBMV	BNYVV	23.72	4.80	1.120
USH-11	BNYVV	BSBMV	18.56	5.33	1.936
USH-11	BNYVV	healthy	9.48	2.03	1.432

Fig. 2



<sup>a samples are listed in order by weight.
b Absorbance values are from the second six week transplant period.</sup>

^c USH-11 typically has a lower yield than B4776R.

Our studies of BNYVV and BSBMV clearly show that both viruses have a significant deleterious effect on the growth of beets. However, whereas BNYVV causes classic Rhizomania symptoms of root necrosis, bearding, reduced sugar and reduced yield, BSBMV does not induce Rhizomania. Far less is known about the effects of BSBMV of sugarbeet crops, except for the fact that reduced yield is expected to occur. Root necrosis is not evident in BSBMV-infected roots in greenhouse experiments using infested soil. Future studies will examine sugarbeet varieties for relative concentrations of BNYVV, BSBMV, and BSBV. We will conduct greenhouse and field trials to estimate effects of these viruses on yield and agronomic traits. Our studies will also attempt to determine if the Rhizomania resistance gene (Rz) to BNYVV has any effect upon the infection and impact of other benyviruses. Through greenhouse and field trials. we will determine if the poor performance experienced in some fields and production regions is due to non-BNYVV benyviruses. We will continue to investigate the existence of other soilborne virus components in the "beet decline" syndromes which have occurred in Nebraska and Colorado. Because Rhizomania and other benyviruses are present in most beet growing regions in the United States, we will conduct greenhouse and field studies to determine the effect of mixed infections on beet performance, and measure their relative levels, compared to single infections. Preliminary evidence suggests that the combination of BNYVV and BSBMV is more severe than either virus alone. This again emphasizes the need for the development of resistance to other soil-borne benyviruses aside from BNYVV.

Western Sugar Company-Grower Joint Research Committee Report. Part I: Investigations into the cause for decreased yield and sugar yield in midwestern sugar beet production.

G.C. Wisler

Introduction

A significant decrease in sugarbeet yield has been observed throughout the Eastern Slopes of the United States for the past few years. Possible causes which have been suggested include Rhizomania, selections of sugarbeet varieties which are not suited to the area in production, or soil-borne fungal, bacterial, and other viral pathogens. Our results suggest that Rhizomania is not the cause, and that other soil-borne sugar beet viruses may have an important role. Our preliminary results indicate that the soil-borne viruses of beet, in particular *Beet soil-borne mosaic virus* (BSBMV) and possibly *Beet soil-borne virus* (BSBV), are important factors in limiting beet production.

The objectives of this study during 1999 were to continue to study the effect of soil-borne viruses that may be associated with the decline. In addition, we conducted a small scale, preliminary field trial to evaluate the effect of fumigation on beet growth, sugar production and the presence of soil-borne viruses.

Materials and Methods:

A. Fumigation Trial: Four varieties were selected for the fumigation trial in Scottsbluff, NE. These were: Monohikari, Beta 4038r, Beta 1399, and Beta 9155. Each variety was replicated six times in both a fumigated section and a non-fumigated section. Four beets were individually dug by hand and topped at the lowest leaf scar at two dates in the growing season; 6-28-99 and 7-9-99. Roots were washed free of soil and scraped for root hairs and epidermal tissue. Samples were transported to Salinas for testing by ELISA for BNYVV and BSBMV, and for mechanical inoculation to indicator plant for miscellaneous viruses. Results from this study are presented in Table 1.

B. Mixed infections: See BSDF 281 report, Table 2.

C. Resistance to BSBMV: See BSDF 281 report, Fig 1.

Results:

Although the fumigation study was not set up in a randomized complete block design due to constraints placed on us which precluded our obtaining statistically significant data, we do see a trend in the fact that fumigation had a positive effect on production as seen by the levels of % sucrose and tons per acre (Table 1). However, it is not clear what is being controlled. There was a very low level of BNYVV and BSBMV as seen in individual plant tests, but this does not show up in our overall averages for these viruses as measured by ELISA. Other than BNYVV and BSBMV, no other virus was detected in this field trial. Further studies are needed in fields known to have a history of infestation by BSBMV in a fumigation trial to measure the true effects of this virus on beet production.

Conclusions:

In several states in the United States both BNYVV and BSBMV have been found. BSBMV has been shown in greenhouse trials to have a significant effect on growth of beets. Resistance to BSBMV has not yet been identified, and the effects of these two distinct viruses appears to be more significant than either one alone, depending on the variety planted. In addition, resistance to Rhizomania does not confer resistance to BSBMV, and efforts should be made to find resistance to this virus. Fumigation in last year's study appeared to show a trend of having a positive effect on beet growth and % sugar. However, this was not an optimum test, for the reason that the field was not heavily infested with BSBMV, and it was not set up in a randomized complete block design. Now that we have identified infested fields, and the crop rotation is back to bees in some fields, we plan to replicate another fumigation test in an infested field. This will depend on the availability of appropriate fields and the cooperation of growers.

Table 1. Comparisons Between Fumigated and Non-fumigated Soil on Sugarbeet Production. Scottsbluff NE. 1999.

		Fun	nigated		
				BNYVV	BSBMV
Variety	% Sucrose	Ton/A	Lb sugar/A	(OD/H)	(OD/H)
Monohikari	14.10	29.66	8368	1.22	1.33
Beta 4038R	12.83	30.86	7916	1.56	0.80
Beta 1399	14.16	25.05	7105	1.23	0.74
Beta 9155	13.61	31.96	8696	1.41	1.03
Mean	13.68	29.38	8021.3	1.36	0.98
LSD (.05)	0.68	4.25	1458.9	0.59	0.52
		Non-F	umigated		
Monohikari	14.00	24.60	6905	1.28	1.10
Beta 4038R	13.32	26.60	7082	1.19	0.86
Beta 1399	14.75	22.50	6655	1.42	1.00
Beta 9155	13.80	29.56	8151	1.33	1.19
Mean	13.97	25.82	7198.2	1.31	1.04
LSD (.05)	0.53	2.46	739.3	0.34	0.66

Part II: Continued study of the new luteovirus causing yellowing in the United States B. Etiology and transmission properties of BChV (collaborators H.-Y. Liu, W. M. Wintermantel, R. T. Lewellen): Since 1995, a yellowing disease of sugarbeet has been frequently observed in Colorado, Nebraska, and California sugarbeet fields. Symptoms of this disease are identical to those caused by beet western yellows virus (BWYV) including interveinal yellowing and necrotic lesions caused by Alternaria sp. BWYV isolates from beet have a wide host range and are readily distinguished by systemic infection of shepherd's purse (Capsella bursa-pastorus) and lack of infection of Chenopodium capitatum. These newly described isolates have a narrow host range and show interveinal reddening on C. capitatum but do not infect shepherd's purse. Biological properties indicate these isolates are distinct from BWYV. This disease is readily transmitted (only one aphid is needed to transmit at an efficiency of 36.6%) (Table 1) in a persistent manner by the green peach aphid (Myzus persicae), but is not mechanically transmissible. The virus has been

purified and the virus particles are isometric and 26 nm in diameter. The coat protein from purified preparations is ca. 23 kDa. This disease may be more damaging to sugarbeet but because of the narrow host range may be more readily controlled by host-free periods than conventional BWYV strains. (This abstract was presented at the 1998 APS meeting, November, 1998.)

Materials and Methods: This year specific antiserum has been developed in our lab which recognizes BChV only. Previously, all antisera to BWYV reacted to both BWYV and BChV, thus hampering a diagnostic test based on serology. Results from our ELISA tests are shown in Table 1.

Serological F	ical Reactions to "Yellows Antisera antisera		
Antigen	BChV	BWYV	
BChV	+	+	
BWYV	-	+	
Healthy	-	-	

Conclusions:

Progress has been made in determining the etiology of BChV its relationship with other yellowing luteoviruses infecting sugarbeet, and sources of resistance to BChV. BChV is now considered to be a new and distinct member of the luteovirus group of aphid transmitted viruses, and a distinct member of the virus yellows complex. Aphid transmission is highly efficient, with only one aphid necessary to transmit BChV. Sources of infection still need to be determined, and specific diagnostic probes to be made available. An ELISA test will be refined this year which is specific only to BChV. Molecular probes will also be made available which recognize each member of the yellows complex individually. Progress is being made in our lab to determine the interactions between the members of the virus yellows complex.

The old and the new: viruses of sugar beet and their impact on beet production in California.

The California Sugar Beet 1999 Annual Report:27-30.

G.C. Wisler

The diverse climate of California lends itself well to a diverse agricultural industry. The variety of weeds, crops, insect and fungal vectors also provide favorable conditions for plant virus disease development. Viruses have had a significant impact on production since sugar beet was first introduced to California, and continue to do so today. Beet curly top curtovirus (BCTV; family Geminiviridae) almost destroyed a fledgling sugar beet industry soon after its establishment in the 1870's. A combination of resistant varieties, cultural and chemical management of beet crops to provide early plant emergence and development, and a highly coordinated beet leafhopper scouting and spray program have allowed for adequate control of BCTV. These programs were initiated by the USDA-ARS in Salinas, California and the University of California, and are still in place today. Populations of the vector of BCTV, the beet leafhopper (Circulifer tennellus) are monitored and can still achieve proportions which can be extremely damaging to the beet crop. Breeding programs continue to evaluate resistance to curly top, as this virus still poses a real threat to production. For example, in 1992 Idaho had the most severe outbreak of curly top in over 20 years, which caused an estimated loss to the sugar beet industry of \$15 million. That was the same year that Rhizomania was first identified in Idaho, thus curly top received relatively little attention. The sugar beet industry should continue to maintain the current scouting program, and breeding programs must continue to improve the resistance that is available for curly top.

"Virus yellows" describes a complex of yellows-inducing viruses that are aphid-transmitted. In the past, this complex has consisted of *Beet yellows virus* (family *Closteroviridae*) and *Beet western yellows virus* (family *Luteoviridae*). *Beet mosaic virus* (BtMV) is often part of this complex, but its importance to the yellowing disease is not completely known. In Europe, *Beet mild yellows virus* (BMYV; family *Luteoviridae*) is part of the virus yellows complex. Early descriptions suggest that virus yellows occurred in the Salinas Valley as early as 1921 when it came to be known as "June Yellows" because by mid-summer. Factors influencing the epidemiology of virus yellows include vector populations, virus/vector relationships and virus sources. From 1950 until the late 1960's, beet yields continuously declined because of increased incidence of the virus yellows complex.

BYV is transmitted in a semi-persistent manner and is retained by the vector for less than 72 hr. This type of transmission suggests that the spread of the virus from the source is local, i.e. the disease incidence is high in areas adjacent to the virus source but quickly decreases with distance. The primary source of BYV is beet itself, including overwintering beets and volunteers in abandoned fields or waste sites. BWYV is transmitted in a persistent manner by aphids, meaning that it circulates through the insect and is maintained for the life of the insect. Thus, distribution of BWYV is more general and widespread than BYV. BWYV infects numerous weeds and other crops, including lettuce. A new component of the yellows complex was identified in the past few years, through the collaborations between the USDA-ARS in Ft. Collins, Colorado and Salinas, and the University of Nebraska. This virus has been identified in Colorado, California, Texas, and in Oregon. It has been named *Beet chlorosis virus* (BChV), a

distinct member of the Family *Luteoviridae*. Symptoms closely resemble BWYV when it is found as a single infection in sugar beet, with intense interveinal yellowing accompanied by *Alternaria* lesions. Symptoms are more orange than yellow in color as with BWYV, and leaves are characteristically thick and brittle. However, these differences are subtle, and BChV and BWYV cannot be differentiated by symptoms alone. Diagnostic tests can include specific nucleic acid tests, serology, and transmission to specific indicator plants by the aphid vectors. The most diagnostic host range difference between these two viruses is that BChV infects *Chenopodium capitatum* but not shepherd's purse (*Capsella bursa-pastorus*), whereas BWYV infects shepherd's purse and not *C. capitatum*. BWYV has a wide host range, whereas BChV has a limited host range. Alternate, or overwintering weed hosts for BChV have still not been identified in areas where BChV has been found. In two years of extensive sampling surrounding weed hosts in Colorado, Nebraska, and California, none was found to be infected with BChV. Thus, the epidemiology of this new virus is not completely known.

Epidemiological studies in the late 1950's by J. E. Duffus established a close correlation between virus yellows incidence and proximity of overwintered beet fields. Sugar beet growers and processors reached agreements to maintain beet-free periods between harvesting and sowing new crops throughout California. This included the destruction of volunteers or "groundkeepers" and weed beets. Because of the diverse planting dates throughout the state due to the diverse climates, beet-free periods differ between beet growing districts. These programs were first introduced in the 1968 crop. Following the introduction of the beet-free period in 1968, the average sugar production in California increased yields by about 40% in the subsequent growing seasons.

Virus yellows re-emerged in 1985 in Northern California as a result of increased aphid populations and erosion of beet-free periods. *Myzus persicae* has been the most common aphid vector of the yellows virus complex. In recent years, however, populations of the black bean aphid, *Aphis fabae*, have increased. Although it is a less efficient vector than the green peach aphid, the black bean aphid has complicated the beet-free periods as a means of disease management because it is more heat tolerant than the green peach aphid and has longer flight periods. This situation extends the period for aphid transmission. Thus, beet-free periods are more important than ever before, and the beet industry has enforced them within beet production districts.

Variety trials conducted in 1997 by R. T. Lewellen of the USDA-ARS, Salinas, showed that the yield reduction caused by BChV was similar but more severe than that caused by BWYV. Sugar yield losses ranged from about 5 to 40%, depending on the variety. The loss pattern for BChV fits the pattern for that of BWYV and BMYV. Lines and hybrids from the virus yellows resistance breeding program at Salinas tended to be the most resistant. The most susceptible commercial hybrids tested were those that have been grown in Colorado and Nebraska where BChV has caused significant damage in the past several years. Future breeding programs in Salinas will continue to evaluate resistance to BChV as a virus yellows complex with BWYV and BYV.

The development of *Lettuce infectious yellows virus* (LIYV) in the southern United States to epidemic proportions, and its apparent disappearance in current cropping systems is an excellent example of the impact that insect populations, cropping patterns and transmission characteristics have on virus ecology and epidemiology. Although LIYV was first described in lettuce, it caused significant losses of sugar beet, cucurbits, and other crops in the southwestern United States for a period of time between 1980 and the early 1990's. LIYV is the type member of the genus *Crinivirus* (family *Closteroviridae*) and is transmitted primarily by the sweetpotato whitefly, *Bemisia tabaci* biotype A. It induces yellowing and necrosis on infected plants

accompanied by a significant reduction in yield. In 1981, lettuce, cucurbits, and sugar beet crops were ubiquitously infected with LIYV, resulting in losses exceeding \$20 million in one growing season. Lettuce yielded 50 to 75% lower than in previous years and sugar beets yielded 20 to 30% less than expected.

Bemisia populations changed during the 1980's and early 90's in the sunbelt states of the U. S., and throughout the tropical and subtropical zones worldwide due to the displacement of biotype A by biotype B. Whereas LIYV is transmitted efficiently by biotype A, it is transmitted 100-fold less efficiently by biotype B. The populations of biotype B increased to astronomical proportions by 1990. The fall melon crops which provided a bridge between consecutive beet and lettuce crops were eliminated due to feeding damage by the B biotype. As a result, LIYV has been virtually eliminated and is no longer found in the southwestern desert.

A second, potentially destructive whitefly-transmitted virus, termed *Lettuce chlorosis virus* (LCV), was found in the Imperial Valley of California after LIYV was no longer present. LCV has several characteristics in common with LIYV, including the typical symptomatology of interveinal yellowing of lower leaves, stiffness of leaves, and leaf necrosis. In contrast to LIYV, both the A and B biotypes of *B. tabaci* are efficient vectors. LCV is similar to LIYV with respect to its host range, except that LCV does not infect members of the *Cucurbitaceae*. The whitefly population is currently being controlled in lettuce by use of the insecticide imidacloprid, and thus LCV has not been an economically significant problem, although it is consistently identified in symptomatic lettuce and sugar beet when yellowing symptoms are found in the southwestern United States. The Imperial Valley sugar beet industry has experienced world record yields in the past few years. Thus, the concern for LCV is low at this time. As resistance to insecticides builds up in the whitefly population, or cropping patterns change, LCV could become a potential epidemiological problem. The industry should continue to monitor for yellowed fields.

Resistance or tolerance to LIYV was developed in lettuce, sugar beet, and cucurbit varieties as a result of the epidemics. Preliminary studies are underway to determine if the resistance to LIYV in lettuce and sugar beet confers resistance to LCV as well. There has been some speculation that perhaps LCV may have been present during the LIYV epidemics and the resistance may be useful for more than one crinivirus infecting these crops. For example, the USDA-ARS sugar beet breeding program for the "virus yellows" complex of luteoviruses and closteroviruses was ongoing in the Imperial Valley when LIYV was prevalent. Because selection was based on absence of yellowing, varieties resistant to LIYV were developed before the causal agent was even characterized.

Rhizomania, caused by *Beet necrotic yellow vein virus* (BNYVV), was first identified in the western hemisphere in 1984 in Paso Robles, California. It has since been identified in Texas, Colorado, Nebraska, Idaho, Minnesota and Oregon. To date, it has not been found in Washington, Michigan, or Ohio. The beet industry is well aware of the symptoms and effect of rhizomania on beet production, and precautions that should be made to prevent its spread. This disease is soil-borne, and is transmitted by the plasmodiophorid fungus, *Polymyxa betae*. The primary source of spread is through the movement of infested soil or beets. Great strides have been made with regard to resistance to rhizomania, and growers are urged to plant these varieties whenever possible in soils known to be infested. In addition, cultural control methods can be employed to manage this disease. These include planting early into cool soils, minimizing overwatering, and crop rotations to reduce levels of virus in the soil. Resistant varieties that are available now yield as well as nonresistant beets under non-rhizomania conditions. Recent studies clearly show that the dose of the *Rz* allele for resistance to rhizomania is directly

correlated with virus levels, rhizomania disease index scores and root weight. With regard to virus levels in infected sugar beets, Rzrz < Rzrzrz < rzrzzrzz.

There are other soil-borne viruses that infect sugar beet in addition to BNYVV. These have been found throughout the United States where beets are grown, with the exception of California. These viruses have not been as thoroughly studied as BNYVV has, thus less is known about the epidemiology of these viruses. In our lab, we have still not detected these "other" viruses from beets grown in California. However, they appear to be fairly widespread in other beet-growing states. Beet soil-borne mosaic virus (BSBMV) is one in particular that has been fairly well characterized. Although every isolate of BSBMV that has been found reacts the same in serological tests, on a molecular level they appear to be a closely related family of viruses, which vary according to their particular ecological niche. This indicates that these virus isolates may actually be endemic to the United States, and have changed and evolved here. BSBMV has not been positively identified in Europe or the United Kingdom. Beet soil horne virus (BSBV) is another soil-borne beet virus which is being found more frequently. In preliminary studies of beets in Nebraska and Colorado, where a significant decline in yield has occurred over the past several years, 24 of 27 affected fields were infested with either BSBMV. BSBV, or both. Only one field was infested with rhizomania. Although these data do not provide conclusive evidence that these viruses are the cause of the declines experienced by growers, the information needs to be investigated further. Years of research has shown that BNYVV causes rhizomania. Other than a low incidence of leaf symptoms for BSBMV that resemble BNYVV leaf symptoms, the virus symptoms for these other soil-borne viruses are not well-known. It is likely, however, that BSBMV and BSBV cause a reduction in yield. How extensive that reduction can be is unknown at this time, and studies are underway to attempt to document that reduction under controlled field conditions. The possible interaction that can occur between BNYVV, BSBMV, and BSBV is another area of concern for researchers and the industry alike. Preliminary studies indicate that two viruses as mixed infections are more severe that either virus alone. Although BSBMV may not exist yet in California, based on our experience with the movement and spread of rhizomania, it is likely that it will be introduced in the future. Pathologists and sugar beet geneticists are preparing themselves for that eventuality by learning more about these viruses and their interactions, and investigating the possibility of resistance to these soil-borne viruses.

Project 220

Viral transgene-mediated resistance to *Beet yellows virus* as a model for engineered virus resistance in sugarbeet

William M. Wintermantel Salinas, California

Introduction: Virus yellows consists of a complex of viruses causing beet leaves to turn yellow prematurely, and has contributed to disease-related losses in California sugarbeet production for many years. This disease complex is composed of members of two main genera of plant viruses, a *Closterovirus* and a *Luteovirus*. Occasionally a *Potyvirus* is also present. Once plants begin showing initial yellowing symptoms, losses accumulate approximately 2 percent each week through the remainder of the growing season. Direct annual losses to virus yellows average in excess of \$36 million, without considering indirect effects such as the displacement of production areas, increased freight costs, and potential loss of processing facilities due to disease-related yield and revenue reductions.

Plant virus resistance obtained through transformation with foreign genes (transgene-mediated resistance) can increase the level of resistance in cultivars which partially control a particular disease, and can provide resistance when none is available through traditional breeding. This project examines the potential for transgene-mediated resistance against Beet vellows virus (BYV) in sugarbeet. BYV is a major component of the virus yellows complex, and has been identified by the California sugarbeet industry as a primary concern. Engineered BYV resistance should complement current resistance/tolerance to Beet western vellows luteovirus (BWYV). the other major viral partner in the virus yellows complex. Transgene-mediated resistance has been studied extensively for a number of years. Since its development in the mid 1980s, transgenemediated resistance has been developed for control of a large number of plant viruses in many different hosts (Baulcombe, 1996; Deom, 1999), including limited attempts to control BNYVV in sugarbeet (Kallerhoff et al., 1990; Ehlers et al., 1991). There are several means by which foreign genes can engender resistance, and often more than one approach can achieve resistance against a particular virus. For example, transgenic resistance has been achieved for tobacco mosaic virus using viral replicase transgenes as well as by using viral coat protein transgenes. The means by which the replicase transgene produces resistance differs from the mechanism by which coat protein-mediated resistance operates, at least for tobacco mosaic virus. The choice of a transgene (the foreign gene being inserted into the sugarbeet genome) must be determined through careful analysis of the interaction between the targeted virus and the sugarbeet plant. The transgene must be able to block the virus infection cycle such that the virus cannot bypass the mechanism of the resistance. It is important, therefore, to have a solid understanding of the nature of the infection process and how disease develops for each virus targeted for transgenemediated resistance. BYV is transmitted by aphids in a semipersistent manner (requiring long feeding times for acquisition and transmission by vectors). In infected plants, BYV is usually restricted to phloem tissues (sieve tubes, companion cells and phloem parenchyma), but is occasionally found in the mesophyll and epidermis near local lesions.

This suggests that strategies which interfere with virus replication and packaging should be effective in generating resistance to BYV.

Purpose: The BYV resistance project is part of a long-term effort to engineer sugarbeets for resistance against plant viruses through transformation with foreign genes, and focuses on development of virus resistance to BYV as a model for using biotechnology to control virus diseases in sugarbeet. The main objectives are as follows:

- 1. Development of nucleic acid constructs for use in plant transformation
- 2. Identification of genes which confer resistance against BYV
- 3. Optimization of sugarbeet transformation and regeneration procedures for select sugarbeet germplasm

Accomplishments: A plant transformation facility was completed in March, 1999, and during the past year we have concentrated on the transformation and regeneration of sugarbeet through tissue culture, as well as development of constructs for sugarbeet transformation with a goal of resistance to beet yellows virus (BYV). Efforts are in progress to improve regeneration efficiency, as this a crucial, limiting step in the sugarbeet transformation procedure. Experiments involving sugarbeet were initiated in the fall of 1999. Constructs for use in plant transformation have been developed, and additional constructs are in progress.

Approach: Cloned BYV genes were generously provided by V. Dolja (Oregon State University, Corvallis, OR). BYV genes have been isolated, modified, and inserted into binary plant transformation vectors. A binary vector, provided by W.R. Belknap (USDA-ARS, Albany, CA) is being used for most transformations to reduce end-product licensing requirements. Plant transformations are being performed using Agrobacterium tumefaciens. Initial transformations are being performed on both sugarbeet and N. benthamiana, an alternate host for BYV. N. benthamiana can be transformed easily using standard procedures (Rogers et al., 1986), and transgenic plants can be tested for resistance to BYV in a fraction of the time required to obtain transgenic sugarbeet. As constructs are identified which provide effective resistance against BYV in N. benthamiana, these will be used for transformation of sugarbeet. Procedures for sugarbeet transformation and regeneration include portions of methods used by Doley and Saunders (1989), D'Halluin et al. (1992), Krens et al., (1996) and others, with some modifications. Transgenic plants will be tested for the presence of the transgene by PCR analysis. Plants exhibiting strong resistance will be subjected to Southern blot analysis to determine the number of copies of the transgene in these plants. A number of different sugarbeet tissues are being used for transformation, including young (not fully expanded) leaf tissue, petiole and bolt tissue. Beet varieties to be used for transformation have been provided by R.T. Lewellen. After resistant, stable transformants are identified, plants will be turned over to R.T. Lewellen for seed production and introduction into the breeding program.

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Project 221

Examination of viral interactions in relation to disease severity and resistance in the virus yellows complex of sugarbeet.

William M. Wintermantel Salinas, California

Introduction: Virus vellows has contributed to disease-related losses in California sugarbeet production for many years. Once plants begin showing symptoms, losses increase approximately 2 percent each week through the remainder of the growing season. This disease complex is composed of members of two main genera of plant viruses, a Closterovirus and a Luteovirus. Occasionally a *Potvvirus* is also present. In California, BYV is the predominant Closterovirus involved, and BWYV and Beet chlorosis virus (BChV) are the principal Luteoviruses in the complex. The Potyvirus, when present, is almost exclusively BtMV. BtMV is generally not considered an economically significant pathogen alone, however BYV, and the Luteoviruses can effect yields substantially in single infections. All of these viruses are transmitted by aphids. particularly the green peach aphid and the black bean aphid. Although beet-free periods are useful in managing virus vellows, the increased range and population of the black bean aphid has made this disease more difficult to control in recent years. Once plants begin showing symptoms, losses increase approximately 2 percent each week through the remainder of the growing season. Traditionally, breeding for resistance has involved breeding for control of the vellowing symptom. BtMV causes symptoms on young plants, but as symptoms of the vellowing viruses develop, mosaic symptoms decrease. Currently, beet varieties are available which exhibit some tolerance to BYV, and field resistance to the *Luteoviruses*. Most commercial varieties do not exhibit substantial levels of resistance to BtMV.

The relationship in sugarbeet between the three viruses in disease induction is not clear. Although all 3 viruses are present in plants at the same time, it was not clear at the beginning of this project whether the yellowing and stunting symptoms associated with the disease are more severe when multiple viruses are present or not. Furthermore, it was not known whether the presence of one virus facilitates or hinders the activity of another. Possible interactions were suggested by observations that yellowing disease and sugar yield reductions were more severe when both BYV and BWYV were present (R. Lewellen, personal communication). It is also noteworthy that BtMV is not considered a serious problem in beet, even though it is often present with BYV and/or BWYV. This study addresses whether synergism or suppression occurs in the virus yellows complex on sugarbeet, and will increase our understanding of the relationship between the virus components of the yellows complex. This knowledge may be helpful in identifying sources of resistance to the yellows complex, and in the development of new resistant varieties.

Approach: Sugarbeet breeding lines have been selected that are either susceptible to, or exhibit a range of resistance levels to each of the three target viruses. These lines are being challenged by aphid-inoculation of BYV, BWYV, and/or BtMV. Plants are inoculated with each virus individually, with all combinations of two viruses, and finally, all 3 viruses are inoculated together. Mock inoculations are also performed with virus-free aphids. Symptom development

is monitored over the course of each experiment, and total nucleic acid samples are prepared from symptomatic leaves. Total nucleic acid concentrations are equilibrated, and replicate dot blots (a form of nucleic acid hybridization) are performed to compare relative levels of each virus in single, double, and triple infections. Relative amounts of viral RNA are compared by phosphorimage analysis of dot blots. At the conclusion of each experiment, soil is removed from roots, and top and root are separated to determine the effects of each virus combination on root and top weight, compared with healthy controls.

Preliminary Results: Sugarbeet containing mixed infections of more than one yellowing virus exhibit greatly increased stunting and reduced root weight compared with single infections and mock-inoculated plants. This pattern is particularly evident in susceptible beet varieties, but also occurs to a lesser degree in varieties exhibiting resistance to BWYV and tolerance to BYV. Effects on beet leaves are not as apparent, particularly when BYV is present. BYV causes a thickening of leaves, resulting in heavier weight.

Experiments in progress are comparing virus titer in sugarbeet infected with each virus individually, mock-inoculated plants, as well as all possible combinations of mixed infections to determine if any synergism or suppression occurs in sugarbeet infected with more than one yellowing virus. Current results suggest substantial synergism between BYV and BtMV, resulting in greatly reduced root weights, but only mild synergism between BYV and BWYV. This data will be correlated with effect of mixed infections stunting severity, root and leaf weight, as well as virus concentration in single and mixed infections.

DEVELOPMENT OF SUGARBEET BREEDING LINES AND GERMPLASM

R.T. LEWELLEN

C26, C27 & C51 - Beta vulgaris L. germplasm lines C26 (PI610488), C27 (PI610489), and C51 (PK593694) were developed by the USDA-ARS in cooperation with the Beet Sugar Development Foundation and the California Beet Growers Association. C26 and C27 were released in 1999. C51 was released in 1996. The germplasm base of sugarbeet (B.vulgaris) is believed to be relatively narrow compared to its ancestral source B.vulgaris L. subsp. maritima (L.) Arcang. Beta vulgaris subsp. maritima (Bvm) occurs from the Mediterranean Basin through the Near East and north along the Atlantic shore to Denmark. The major geographic ecotypes of Bvm are the easy bolting annuals of the Mediterranean Basin and the harder bolting annuals or biennials of western Europe. Both of these ecotypes are difficult to evaluate directly for reaction to diseases and pests and for agronomic traits. Even though fully cross compatible with sugarbeet, synchronizing flowering of Bvm with biennial sugarbeet is often difficult. To overcome these problems and to make the broadly based germplasm of Bvm more accessible for evaluation and breeding, groups of Bvm accessions were crossed to sugarbeet and improved. The broadly based germplasms C26, C27, and C51 are populations from these prebreeding programs at Salinas.

C26 is a multigerm, self-fertile line that theoretically received 50% of its germplasm from sugarbeet and 50% from B. vulgaris subsp. maritima (Bvm). The wild sea beet Bvm principally was derived from accessions collected by Doney et al. in France, UK, and Ireland. C26 was developed from crosses between sugarbeet line C37 and Bvm. The sources of the Bvm plants were from accessions tested in the 1991 and 1993 Sugarbeet Crop Germplasm Committee (CGC) sponsored tests at Salinas. Plants from within individual PI accessions that showed high resistance to rhizomania caused by beet necrotic yellow vein virus were selected. In 1991, about 200 selected plants from 20 accessions collected in the UK and 6 accessions collected in Ireland were bulked and increased in mass in 1992 to produce a Bvm population called R423 (PI599350). In 1993, about 160 rhizomania resistant plants from 11 PI lines collected in France were bulked. Stecklings from population R423 and the bulked selected plants from the French accessions were combined into a single pollinator in 1994 and crossed in bulk to C37. Seed from the Bvm plants were called R423B (PI599351). C37 is uniformly susceptible to rhizomania and has only green hypocotyls. Seed harvested from the C37 seed-bearing plants was sown in August 1994 into a field plot with rhizomania infestation. In December 1994, F₁ plants were selected and increased in 1995 by open pollination to produce an F₂ population called R526. Records were not maintained as to the contribution of each wild beet accession or which accessions were specifically involved. The UK accessions were in the PI518298-518372 (WB620-694) series. The Irish accessions were in the PI518381-518416 (WB703-738) series. The French accessions were in the PI540598-540608 (WB852-862) series.

Plants from the F₂R526 population were grown in the field under rhizomania infested conditions and were inoculated with beet yellows and beet western yellows viruses, the cause of virus yellows (VY) *Erwinia carotovora* (Jones) Bergey et al. subsp. *betavasculorum* Thomson et

al. and *Erysiphe polygoni* DC, the cause of powdery mildew. Individual plants were selected for resistance to rhizomania, Erwinia root rot, powdery mildew, and for nonbolting, beet conformation, root size, and sucrose concentration. Selection for VY was indirect and was based upon individual root performance for root and sugar yield. Selected roots were increased in mass by open pollination to produce F₃ line R726. From R726, two successive selections were made under field conditions for resistance to rhizomania, nonbolting, and root conformation and size and increased to produce the F₅ line R926 that was released as C26.

C26 should have performance and traits similar to its F₃ and F₄ generations. The F₃ line R726 and F₄ line R826 have been evaluated in field trials at Salinas and Brawley, CA. They have shown high resistance to rhizomania. Most plants appear to be biennial or hard bolting annuals. Pigmentation is mostly similar to that of sugarbeet but some *Bvm* patterns still occur. Under rhizomania and/or VY conditions, the components of yield except for lower juice purity are similar to other open-pollinated lines of sugarbeet. Under VY infected conditions, R726 has yellowing symptoms that score similar to the most tolerant sugarbeet lines. Under mild cercospora leaf spot epiphytotic at Salinas, R726 was moderately resistant but in subsequent tests in CO and MN, it was moderately susceptible. C26 has a dark green canopy, similar to the coloration of many *Bvm* lines from NW Europe but without the thickened leaves. C26 is an enhanced, broadly based population from which useful genetic variability might be found for the future improvement of sugarbeet.

C27 is a multigerm, self-sterile line with sugarbeet and B. vulgaris subsp. maritima (Bvm) germplasm. The sea beet Bvm component was selected from accessions being evaluated for resistance to rhizomania in the 1996 Sugarbeet CGC sponsored test at Salinas. About 200 Bvm rhizomania resistant plants from 19 accessions were selected. These accessions represented introductions from UK (PI518426, PI518435, and PI518440), Poland (PI535833, PI535835, and PI535843), and France (PI540568, PI540575, PI540588, PI540593, PI540596, PI540598, PI650599, PI540600, PI540601, PI540602, PI540603, PI540604, and PI540605). After vernalization, the selected Bym plants were bulked and transplanted into a seed isolation plot with sugarbeet lines C37 and C69. All plants in the seed plot could have crossed inter se. Seed from the Bvm plants was called R720 (PI599352). Seed harvested from C37 was called R727A and that from C69 was called R727B. These seed lots would have contained sibmated, interline crosses, and F₁ individuals. Resistance to rhizomania and wild beet plant type and color patterns were used to identify sugarbeet x Bvm F₁ plants. Selected F₁ plants from both sugarbeet parents were bulked and increased to produce a single F₂ population called R827. From R827, beets were selected for resistance to rhizomania, nonbolting, root size, and beet conformation and increased to produce the F₃ population R927 that was released as C27.

C27 segregates for high resistance to rhizomania. Resistance could have been derived from C69 (factor Rz) and/or Bvm. The allelism or uniqueness of the resistance from Bvm to Rz and other previously identified sources of resistance is not known. C27 has had limited agronomic evaluations but should be broadly based, enhanced germplasm from which new genetic variability can be identified for the future improvement of sugarbeet.

C51 is a self-sterile, multigerm, germplasm line that theoretically received 50% of its germplasm from sugarbeet and 50% from *B.vulgaris* subsp. *maritima* (*Bvm*). The *Bvm* germplasm was derived from a collection of about 60 accessions collected primarily from the Mediterranean Basin. C51 is an advanced version of C50 (PI564243 and PI59079) that has been improved for sugarbeet traits and disease resistance. From C50 [=F₃ (sugarbeet line C54 x *B.vulgaris* subsp. *maritima*)], improved subpopulations were created by four to six cycles of

recurrent phenotypic selection for various combinations of productivity and host-plant resistance. Selections were made for biennialism, root and crown conformation, sucrose concentration, and root yield. Concurrently, selections were made for resistance to rhizomania, and/or VY. In 1995, mother roots selected for sucrose concentration and yield under severe rhizomania conditions from eight or these subpopulations were recombined to for C51. The component lines of C51 have been tested as versions of breeding line R22, e.g., R422Y3 and R422R5. C51 was released and evaluated as breeding line R522.

Subpopulation components of C51 (R22R lines) that had been selected for resistance to rhizomania have performed very well under severe rhizomania conditions. In tests at Salinas and Brawley, CA, they often have had comparable sugar yield to commercially available rhizomania resistant hybrids. At Brawley under rhizomania conditions, these lines have shown the best known resistance to high temperature root rots and plant death. There is evidence that a factor or factors in C51 conditions a higher level of resistance to rhizomania (BNYVV) than that conditioned by Rz, the Holly gene. Experimental hybrids show that this factor in C51 is expressed in a dominant manner.

Subpopulations of C51 (R22Y lines) that had been selected for VY resistance on the basis of sugar concentration and yield have performed relatively well under both VY infected and noninfected conditions as compared to normal VY tolerant sugarbeet lines. Under nondiseased conditions, these R22Y lines have shown surprisingly high sucrose levels. These results suggest that C51 might be a source for new genetic variability for sugar concentration and yield as well as disease resistance.

C51 likely will be most useful in the near term as a source for high levels of resistance to rhizomania and for plant persistence under the combined effect of rhizomania and high temperature conditions. Resistance to rhizomania from C51 has been backcrossed into C37 and released as C79-8 and into other sugarbeet backgrounds and released as C67 (PI599340) C72 (PI599342) and C890-8. In a longer term, C51 should provide useful genetic variability for resistance or tolerance to virus yellows, other sugarbeet diseases and pests and possible for components of sugar yield productivity.

DOWNY MILDEW - Downy mildew caused by *Peronospora farinosa* had a high incidence in the November planted bolting evaluation trials (Tests 199-999). Prior to attempted control with Ridomil-Gold MZ, counts of visibly infected plants were made on May 26, 1999. In the late 1940s and 1950s, resistance to downy mildew (DM) was one of the main breeding objectives at Salinas. DM can be severe in winter planted beets in the coastal states including the seed fields of Oregon. Since about 1965, selection for resistance to DM has been a minor part of the breeding program. It appears that in this time, there has been a shift from a moderately resistant germplasm base to a more susceptible base. However, the older lines known to have been resistant in the 1960s still appear to be moderately resistant, e.g., C562. Considerable variability among and within breeding lines for reaction to DM is evident in these 1999 tests. For example, C76-89-5 appears to be one of the more resistant breeding lines (Test 299, 199, 499). In Test 499, full-sib lines generated from C76-89-5 to evaluate for nonbolting. components of yield, and resistance to diseases (e.g. virus yellows, rhizomania, Erwinia, powdery mildew) were evaluated for reaction to DM. Individual FS's from C76-89-5 ranged from 0 to 26% infected plants whereas, FS's from other lines showed up to 92% infection. In test 699, S₁'s from the F₁ hybrid of C76-89-5 x popn-931 ranged from 0 to 100% infected. S₁lines

from popn-931 (Test 799) ranged from 2 to 86% infected. These counts suggest a significant genetic component for reaction to DM. At some point in the future, it may be of interest to do a genetic analysis of this resistance. To my knowledge the inheritance of downy mildew resistance in sugarbeet has never been elucidated.

INDEX OF VARIETY TRIALS, SALINAS, CA, 1999 U.S. AGRICULTURAL RESEARCH STATION

Tests were located in three field plot areas at Salinas and two at Brawley, CA. Disease nurseries were also used in Idaho, Colorado, and Minnesota. Tests at Brawley (Imperial Valley) were planted in September and October 1998, and harvested from May through July, 1999. Tests at Salinas were planted from November, 1998, through August, 1999, and harvested from September through December. Tests at Spence Field (Salinas) were under both rhizomania and nonrhizomania (following methyl bromide fumigation) conditions. Herbicides were not used in Block 6 trials that followed strawberries and methyl bromide fumigation. Nortron, Pyramin, Betamix, Progress, and Poast were used in the other trials. Bayleton at 2lbs material/acre was used for powdery mildew control. Lorsban-4E was applied for aphid and other insect control. The specific planting and harvest dates as well as plot size and design are shown on each test summary.

Tests are listed in the main Table of Contents for Salinas by types of material and evaluation. As an aid to find test summaries, they are listed below by ascending test (planting date) number and cross-referenced to the page number. Tests shown as N/A are not available or included in this report.

TEST NO.	NO. ENTRIES	TEST DESCRIPTION	PAGE NO.
<u>110.</u>	ENTRIES	TEST DESCRIPTION	<u>110.</u>
PROGEN	Y TESTS FO	R NONBOLTING, VIRUS YELLOWS & RHIZOMANI	<u>A</u>
100 1	4.0	T	A 1 (5
199 et al.	48	Testcross hybrids of selected S ₁ MM lines	A165
399 et al.	80	Full-sib progeny from lines with <i>Bvm</i> gp.	A171
499 et al.	112	Full-sib progeny fromC31 <i>Rz</i> -type lines	A178
599 et al.	64	Full-sib progeny from C78 & C80	A184
699 et al.	80	S_1 progeny from F_1 (MM, S^1 , Aa, Rz) lines	A188
799 et al.	48	S ₁ progeny from MM.S ^f .Aa,Rz populations	A193
899 et al.	128	S ₁ progeny from mm, S ^f . Aa. Rz populations	A196
999 et al.	72	Topcross hybrids of S ₁ mm progeny lines	A204
BOLTIN	G EVALUATI	ON TEST, BLOCK 2S, PLANTED, NOV. 1998	
199	80	Nonbolting evaluation of hybrids	A156
299	160	Nonbolting evaluation of breeding lines	A149
999	96	Nonbolting evaluation of topcross hybrids	A160

TEST NO.	NO. ENTRIES	TEST DESCRIPTION	PAGE NO.
VIRUS	YELLOWS (BY	V-BWYV-BChV) PROGENY TESTS, BLOCK 3,	
	ED APRIL 1999	······································	
1100	20		
1199	32	VY evaluation of selected MM S ₁ progeny lines	n/a
1299	64	VY evaluation of lines (BTS)	n/a
1399	48	Full-sib progeny from lines with <i>Bvm</i> gp	n/a
1499	112	Full-sib progeny from C31Rz-type lines	n/a
1599	64	Full-sib progeny from C78 & C80	n/a
1699	80	S ₁ progeny from F ₁ (MM.S ^f ,Aa.Rz) lines	n/a
1799	32	S_1 progeny from MM. S_1^f .Aa, Rz populations	n/a
1899	48	S ₁ progeny from mm,S ^f ,Aa,Rz populations	n/a
<u>VIRUS</u>	YELLOWS (BY	V-BWYV-BChV) EVAL., BLOCK 5, PLANTED MA	ARCH 1999
2000	24	NV	4.52
2099	24	VY evaluation of topcross hybrids	A 52
2199	48	VY evaluation of breeding lines	A 36
2299	48	VY evaluation of experimental hybrids	A 49
	RUS YELLOW ED MARCH, 19	'S INOCULATED COMPANION TESTS, BLOCK 5	
2399	72	Evaluation of topcross hybrids	A 64
2499	48	Evaluation of breeding lines	A 39
2599	48	Evaluation of experimental hybrids	A 54
2377	10	Evaluation of emperimental by entag	
YIELD'	TRIALS, BLOC	CK 5, PLANTED MARCH, 1999	
2699	48	Evaluation of experimental hybrids	A 57
2799	24	Evaluation of topcross hybrids	A 62
2899	24	Evaluation of population hybrids	A 60
2999	24	Evaluation of monogerm lines & populations	A 45
ERWIN 1999	IA ROOT ROT	/POWDERY MILDEW EVAL., BLOCK 3, PLANTI	ED APRIL
3199-2	21	Evaluation of powdery mildew (USDA entries)	A139
3199-2 3299	40	CBGA coded powdery mildew (OSDA entries)	n/a
	168	Inheritance of PM resistance	n/a
3399			A140
3499	80	ERR/PM eval. of MM breeding lines	A140 A144
3599	40	ERR/PM eval. of progeny lines	A144 A146
3699	40	ERR/PM eval. of mm populations	A140

TEST NO.	NO. ENTRIES	TEST DESCRIPTION	PAGE NO.
YIELD'	TRIALS UNDE	R RHIZOMANIA, BLOCK 2S, PLANTED APRIL 1999	
3999	49	Observation & BNYVV titer	n/a
4099	12	Observation & selection (Seedex)	n/a
4199	12	Observation of Nematode resistance	n/a
4299	60	Evaluation of Pl's and Salinas lines	n/a
4399	80	Full-sib progeny from lines with Bvm gp	n/a
4499	112	Full-sib progeny from C31Rz-type lines	n/a
4599	64	Full-sib progeny from C78 & C80	n/a
4699	80	S ₁ progeny from F ₁ (MM.S ¹ .Aa.Rz) lines	n/a
4799	48	S ₁ progeny from MM.S ¹ .Aa,Rz populations	n/a
4899	128	S ₁ progeny from mm,S ¹ .Aa.Rz populations	n/a
4999	24	Performance of populations & lines	A 47
5099	48	WS.BTS.USDA hybrid evaluation (RI-IV)	A 80
5199	78	CBGA coded evaluation (RI-IV)	A 84
5299	18	Population hybrids	n/a
5399	48	Evaluation of MM lines & populations	A 42
5499	48	Evaluation of testcross hybrids	A 68
5599	72	Evaluation of topcross hybrids	A 76
5699	24	Evaluation of population hybrids	A 74
5799	9	Mother root selection	n/a
5899	48	Evaluation of experimental hybrids	A 71
5999	96	S ₂ progeny evaluation	n/a
6099	144	Progeny evaluation for homozygosity	n/a
SELECT	TION FOR RES	SISTANCE TO RHIZOMANIA & POWDERY MILDEW	,
	2M, AUGUST 1		2
6199	57	1999 MM seed productions from gh & isolators	n/a
6299	28	1999 mm seed productions from gh & isolations	n/a
6399	15	1999 seed productions from field isolations	n/a
6499-1	24	1999 increases of selected mm S ₁ progeny	n/a
6499-2	48	1999 increases of selected MM S ₁ progeny	n/a
6499-3	48	1999 increases of nematode resistant lines	n/a
6499-4	144	S ₁ progeny being O-type indexed	n/a

TEST NO.	NO. <u>ENTRIES</u>	TEST DESCRIPTION	PAGE NO.
<u>IMPERI</u>	AL VALLEY		
NONRH	IZOMANIA YI	ELD, FIELD J, PLANTED SEPTEMBER, 1998	
B199	32	Evaluation of testcross hybrids	A 89
B299	32	Area 5 coded variety trial	A 95
B399	32	Evaluation of experimental hybrids	A 91
B499	16	Evaluation of topcross hybrids	A 93
RHIZON	MANIA YIELD	(MILD), FIELD K, PLANTED SEPT/OCT., 1998	
B599	32	Area 5 coded rhizomania	A110
B699	48	Evaluation of experimental hybrids	A 99
B799	72	Evaluation of mm S ₁ progeny topcrosses	A102
B899	72	Evaluation of MM S ₁ progeny testerosses	A106
		VATION (SEVERE DISEASE), FIELD K, PLANTED YALUATED JULY, 1999	
B999		Early evaluation for rhizomania	n/a
B1099	32	Evaluation of testcross hybrids	A114
B1199	64	Evaluation of MM breeding lines	A116
B1299	128	Full-sib & S ₁ evaluation for survival	A120
B1399	138	S ₁ evaluation of mm lines for survival	A126
TRANSO	GENIC HYBRII	D EVALUATION, FIELD J, OCTOBER, 1999	
B1499	6	Evaluation of herbicide transgenics	A132
BSDF C	URLY TOP NU	RSERY, KIMBERLY IDAHO, 1999	
USDA	180	Beet curly top evaluation	A134
CERCO	SPORA LEAF S	SPOT EVALUATION	
USDA	20	CR evaluation at Ft. Collins, Shakopee & Italy	A148
		. 1	

PERFORMANCE OF LINES UNDER VIRUS YELLOWS INFECTION, 1999 TEST 2199.

48 entries x 8 reps, RCB(E) 1-row plots, 21 ft. long

Inoc. BYV/BWYV/BChV: June 7, 1999 Harvested: October 5-7, 1999 Planted: March 22, 1999

		Ac	Acre Yield	7		Beets/					
Variety	Description	Sugar	Loss	Beets	Sucrose	100,	RJAP	Vi	Virus Ye	Yellows	
		Tps	ઋા	Tons	& 	No.	∞	07/21	08/04	08/24	Mean
2199-1: M№	MM, O. P. lines										
B4035R	Betaseed, 7-10-97	8567	32.0	27.20	15.74	162	83.9	4.5	4.9		5.2
KW6770	Betaseed, 6770.5193,1-10-97	7419	43.5	21.65	17.14	2	82.8	5.4	5.9	7.3	6.0
97-SP22-0	Inc. SP7622-0	3962	61.3	٠.	9.	159	80.3	5.9		-	6.5
98-EL-04/02	2 RZM(C80 x EL-smooth root)	7247	41.9	ω.	15.24	9	83.6	•	5.1	9.9	5.4
R876-89-5NB	B RZM-8S R576-89-5NB	8596	24.8	6.2	16.44	168	84.7	2.9	3.4	4.3	3.3
R881	RZM R776,R781,R781-43,	10102	9	2	15.66	Ŋ	83.9	3.3		5.4	4.0
R882	Inc. R776,R781,R781-43,	6686	24.5	30.50	15.40	155	84.7	•	3.6	5.0	4.0
R878%	RZM R778%	9072	29.5	æ.		S	83.6	3.8	3.9	5.5	4.3
R880	RZM R780	9318	28.4	9.4	15.81	5	83.9		4.1	•	4.5
Y868	RZM Y768	9755	25.5	9.9	7	161	84.3	3.8	•	5.3	•
X869	RZM Y769, (C69)	9975	ij.	31.05	16.09	146	83.7	3.1	3.1	•	3.7
X871	RZM Y771	9284	31.3	0.5	15.25	161	83.4	3.8	•	5.8	4.3
Y872	RZM-8S Y672	9233	26.7	30.25	15.26	162	81.9	3.8	4.3	6.0	4.5
Y872B	RZM Y772, (C72)	0006	28.0	9.2	15.44	9	81.9	4.0	4.6	0.9	4.8
X875	RZM Y775	9419	24.8	•	9.	2	81.8	3.3	3.6	5.3	•
Y875 (Sp)	RZM Y775,Y773,Y772,Y767	9324	28.5	0.0	15.55	2	82.9	•	4.1	•	4.3
Mean		8729.5	l. 1	27.80	15.67	157.4	83.4	4.0	4.2	5.8	4.6
LSD (.05)		756.8	ļ. 	2.35	0.42	11.8	٠	0.5	0.5	•	0.4
C.V. (%)		8.8	 - 	8.55	2.69	7.6	2.1	13.8	•	9.6	8.5
F value		30.6	1.1**	28.77**	24.46**	1.7N	s 4.	19.2**	22.0**	24.4**	38.2**
TEST 2199.	PERFORMANCE OF LINES UNDER VIRUS YELLOWS	/IRUS YELL		INFECTION, 199	66						

0.6 0.4 -., 12.7 10.1 8.5 11.1** 15.6** 17.8**27.3**

6.0 0.6 10.1

4.3 0.5 12.7

3.9 0.6 14.7

82.8 1.6 2.0 3.9**

158.4 12.6 8.0 1.2NS

15.45 0.45 2.95

8490.2 27.42 828.8 2.55 9.9 9.43

LSD (.05) C.V. (%) F value

Mean

18.7**15.88** 17.36**

48 entries x 8 reps, RCB(E). ANOVA to compare means across sets.

(cont.)

			Ą	Acre Yield	ק		Beets/					
	Variety	y	Sugar	Loss	Beets	Sucrose	1001	RJAP	ίV	Virus Ye	Yellows	Ì
			rps	%	Tons	ον	No.	o⊱	07/21	08/04	08/24	Mean
	2199-2:	MM lines with WB germplasm										
	SS-432R	Spreckels, 2-8-99	9442	7.	9.5	6.0	9	2	•	•	•	•
	B4776R	Betaseed 4776R, 1-19-99	8740	39.4	26.55	16.46	161	84.5	5.0	5.9	7.8	6.0
	97-US75	Inc. 268 (US75) susc.ck	5337	7.	0.0	3.3	9	0	•	•	•	•
	97-C37	Inc. U86-37	7191	0	3.7	5.1	Ŋ	2	•	•	•	•
	P813	Inc. 6201-#,6202-#s(C),(CP01)	6937	ζ.	3.0	5.0	9	2	•	4.1		4.3
	P814	Inc. 6205-#, 6206-#s(C), (CP02)	7227	26.1		15.26	159	82.0	3.1	3.5	5.5 4	Н
у З	R879	RZM R779 (C79-1, Rz)	6380	2	2.3	4.3	4	8	•	4.5	•	4.8
7	R836	RZM R736, R746 (C79-8,R22)	6982	H.	4.0	4.5	9	•	•	•	6.9	4.7
	R853	RZM-ER-%S R653, (BC4)	6611		1.9	5.0	2	2	•	•		•
	R854	RZM R754, (BC ₅)	7890	27.2	6.1		9	83.2	•	4.1		•
	X873		8506		26.80	ω.	158	8	3.6	•	6.3	4.5
	X873B	RZM Y773	8052	28.2	7.1	14.84	9	82.1	•	4.4	•	4.7
	R840	RZM R740 (C79-#s)	8069	•	7.2	4.8	2	2.	•	9.6	•	4.7
	P811	RZM-PMR 6203-6208-#(C)	8003	31.8	6.8	4.9	9	2	•	•	•	
	X866	RZM Y766	8611	30.0	27.25	15.83	151	83.2	4.1	4 . 4	5.6	•
	X867	RZM Y767, (C67)	9253	25.9	9.4		154	e.	•	•	•	4.0
	Mean		7701.9	1.	ω.	Η.	158.9	82.4	•	4.2	•	4.7
	LSD (.05)	(0	978.8	1. 1	5.35		급.	Η.	9.0	•	9.0	0.4
	C.V. (%)		12.8	ļ. 1	ㄷ.	3.22	7.2	1.9	•	13.9	9.5	9.8
	F value		10.13	1.1 **	6.30**	19.09**	•	w w	5.9**	11.3**	14.6**	16.7**

PERFORMANCE OF LINES UNDER VIRUS YELLOWS INFECTION, 1999 TEST 2199.

(cont.)

		Ac	Acre Yield	ט		Beets/					
Variety	Description	Sugar	Loss	Beets	Sucrose	100'	RJAP	Λ	Virus Yellows	llows	
		Irbs	ov∘ I	Tons	. %	No.	∞ા	07/21	08/04	08/24	Mean
2199-3: N	MM, Sf, Aa populations										
B4419R	Betaseed, 1-19-99	9284	32.1	8.8	6.0	165	4.		5.3	7.1	5.6
Rifle	Spreckels, 2-8-99	8104	41.8	5.4	5.9	156	щ	5.1	6.1	7.6	•
8931	RZM 7931,6915,6925(C)aa x A	10635	18.7	34.35	15.49	158	82.5	3.0	3.6	4.6	3.8
2831	RZM Z731,Z730,Z725(C)aa x A	8769	36.9	7.8	5.7	162	m.	4.8	4.9	6.3	•
8924	RZM 7924, aa x A	9337		ω.	15.64	156	83.1	9.6	4.4	9.	4.7
8926 (Sp)	7931aa x RZM 7926	10256	19.9	ω.		Ŋ	ю	3.3		•	•
	RZM 7926, aa x A	9591	24.9	. 7	15.63	160	8	3.6	3.9	5.5	4.3
8932M	7932CT, 7201-7215Maa x A	9035	25.9	œ ·	15.64	Ω	83.1	3.8	•	•	•
P812	RZM-PMR 6211-#-6217-#(C)	9032	о О	9.6	15.24	155	82.1	3.5	3.4	4.6	3.7
CR811	RZM CR711, (CR09/10)	8319	36.6	4.	15.16	155	81.9	4.0	4.5	6.5	•
CR812	RZM CR712	8347	H	ė.	5.5	5	2	4.3	4.8	•	5.2
CR813	RZM CR713	7944	35.5	٦.	14.69	163	81.2	4.1	4.6	9.9	5.0
N730	Inc. N629, N630 (galls)	7994	30.9	6.6	15.00	159	82.6	4.0	4.6	5.8	4.7
8935	RZM R776-89-5H13	8607	ė.	7.6	15.56		81.4	•	3.4	5.0	3.7
8936	RZM R776-89-5H31	10036	19.8	31.30	16.04	161	83.4	3.1	3.6	5.0	3.8
8939	RZM Y769H31	9337	9	9.9	15.60		82.9	•	4.0	5.4	4.3
Mean			1. I	29.12	15.52	159.0	82.7	•	4.3	•	4.6
LSD (.05)		719.5	l .	2.22	0.42	9.6	1.6	9.0	0.5	9.0	0.4
C.V. (%)		8.0	l. I	7.69	2.72	6.1	٠	٠	12.0	10.5	8.2
F value		10.1*	1. 1	9.62**	6.19**	1.0N	1S 2.0*	11.0**	16.7**	17.0**	31.5**

Notes: Test 2199 was inoculated with virus yellows (BYV-BWYV-BChV) on June 7, 1999. This is two weeks earlier than tests 2099 and 2299 were inoculated. This helps account for the lower yields and greater estimated yield loss of this Relative % loss values were calculated from non-VY test 2499. test.

TEST 2499. PERFORMANCE OF LINES, SALINAS, CA., 1999

48 entries x 8 reps, RCB(E) 1-row plots, 21 ft. long

Planted: March 24, 1999 Harvested: September 29, 1999

		,	7				
		Acre Yleld	eld		Beets/		
Variety	Description	Sugar	Beets	Sucrose	100,	Bolting	RJAP
i		The	Tons	o\ °	No.	%	%
2499-1: MM,0.	MM,O.P. lines						
B4035R	Betaseed, 7-10-97	13598	0.0	ė.	155	0.4	84.7
KW6770	Betaseed, 6770.5193, 1-10-97	13124	5.6	8	152	0.0	5
97-SP22-0	Inc. SP7622-0	10240	33.45	15.34	161	0.0	85.5
98-EL-04/02	RZM (C80 x EL-smooth root)	12483	6.6	5	156	0.0	4.
R876-89-5NB	RZM-%S R576-89-5NB (C76-89-5)	142	w.	7	165		~
R881	RZM R776, R781, R781-43,, (C82)	12024	ω.	S	152	0.0	85.8
R882		12446	39.20	15.81	158		$^{\circ}$
R878%	RZM R778%, (C78)	12873	38.53	9	150	0.0	84.3
R880	RZM R780, (C90)	13019	•	6.2	158	0.0	82.1
X868	RZM Y768	13088	39.43	16.59	153	•	86.1
X869	RZM Y769, (C69)	99	•	6.4	141	•	84.0
Y871	RZM Y771	13509	42.95	15.73	158	0.0	83.8
Y872	RZM-%S Y672	20	9.3	6.0	9	•	
X872B	RZM Y772, (C72)	50	9.0	٥.	2	•	m
X875	RZM Y775	12532	38.10	16.41	159	0.0	83.7
Y875 (Sp)	RZM Y775, Y773, Y772, Y767	04	0.0	6.3	Ŋ	•	ω.
Mean		12573.1	38.51	16.33	155.7	0.02	83.9
LSD (.05)		1117.0	2.84	0.75	•	0.25	1.9
C.V. (%)		0.6	7.46	4 . 62	6.0	1147.90	2.3
F value		4.2**	5.75**	7.76**	3.0**	1.00NS	•
	SEORMANCE OF LINES, SALINAS,	66					
ntries x	8 reps, RCB(E). ANOVA across tests	to compare	means.				
Mean		211.	37.67	16.19	ر. كا	0.91	83.8
LSD (.05)		1073.5	2.89	9.	11.9	1.89	2.0
C.V. (%)		თ თ.	7.79	7	7.7	209.90	2.4
F value		4*1.6	7.69**	7.75**	1.5*	24.70**	2.1**

TEST 2499. PERFORMANCE OF LINES, SALINAS, CA., 1999

(cont.)

		Acre Yield	ield		Beets/		
Variety	y Description	Sugar	Beets	Sucrose	1001	Bolting	RJAP
		Ibs	Tons	o⁄o	No.	ov	o/o I
2499-2:	MM lines with Bvm germplasm						
	Spreckels, 2-8-99	13029	38.90	16.74	159	0.0	84.1
B4776R	Betaseed 4776R. 1-19-99	14423	41.20	17.49	162	0.0	85.9
97-US75	Inc. 268 (US75) susc. ck	10183	34.25	14.89	162	•	83.5
97-C37	Inc. U86-37	10288	32.45	15.85	158	0.0	83.2
P813	Inc. 6201-#, 6202-#s(C), (CP01)	10214	33.05	15.45	161	15.6	84.1
P814	Inc. 6205-#, 6206-#s(C), (CP02)	9781	31.10	15.73	154	9.1	81.6
R879		9486	32.60	14.56	147	0.0	83.1
R836	RZM R736, R746 (C79-8, R22)	10163	32.85	15.48	168	0.4	
R853	RZM-ER-%S R653, (BC4)	10539	34.40	15.32	167	0.0	85.5
R854	RZM R754, (BC ₅)	10832	34.95	15.43	158	0.0	85.2
X873	RZM-ER-%S Y673	12403	38.00	16.31	162	0.0	83.4
Y873B	RZM Y773	11217	36.25	15.48	158	0.0	84.0
R840	RZM R740 (C79-#s)	11059	34.85	15.89	155	0.8	83.6
P811	RZM-PMR 6203-6208-#(C)	11737	7.0		163	15.6	83.2
X866	RZM Y766	12469	37.45	16.64	146	0.0	83.8
X867	RZM Y767, (C67)	12495	37.90	16.49	151	0.0	84.1
Mean		11269.8	35.45	15.85	158.2	2.6	83.8
LSD (.05)		1015.0	2.69	0.72	8.5	3.3	•
C.V. (%)		9.1	7.68	4.57	5.4	127.4	2.5
F value		14.3**	8.37**	8.34**	4.3**	22.7**	•

TEST 2499. PERFORMANCE OF LINES, SALINAS, CA., 1999

(cont.)

Variety	Description	Acre Y. Sugar	Yield Beets	Sucrose	Beets/ 100'	Bolting	RJAP
		I.bs	Tons	o≯e	No.	ok∙	oڥ
2499-3: MM, S B4419R	S ^f , Aa populations Betaseed, 1-19-99	13663	40.20	17.01	159	0.0	85.4
Rifle		393	40.60	17.20	2	0.0	
8931		13089	41.05	15.94		•	83.4
2831	RZM Z731, Z730, Z725(C)aa x A	13887	41.70	16.66	153	0.0	84.1
8924	RZM 7924, aa x A	13512	41.30	16.35	154	0.0	84.4
8926 (Sp)	7931aa x RZM 7926	12807	38.91	16.48	152	0.0	82.5
8927	RZM 7926, aa x A	12779	40.24	15.86	149	6.0	82.4
8932M	7932CT, 7201-7215Maa x A	12189	36.65	16.63	158	0.0	83.9
P812	RZM-PMR 6211-# - 6217-#(C)	12809	40.13	15.95	158	0.4	83.5
CR811	RZM CR711, (CR09/10)	13130	40.70	16.16	2	0.0	83.9
CR812	RZM CR712	12159	37.15	16.40	154	0.0	84.1
CR813	RZM CR713	12325	38.50	16.00	156	0.0	84.7
N730	Inc. N629, N630 (galls)	11576	5.8	16.15	152	0.8	83.0
8935	RZM R776-89-5H13	11630	34.98	16.64	152	0.0	82.4
8936	RZM R776-89-5H31	12518	œ	16.46	154	٠	84.1
8939	RZM Y769H31	12650	38.90	16.27	155	0.0	83.4
Mean			0.	16.39	154.0	0.1	83.7
LSD (.05)		1053.8	3.14	0.53	10.4	9.0	2.0
C.V. (%)		•	۲.	3.26	6.8	482.2	2.4
F value		3.7**	3.32**	4.11**	0.5NS	1.8*	1.4NS

rhizomania conditions. Test 2499 was produced under what appeared to be nearly disease free conditions in soil previously fumigated with methyl bromide for strawberry production. Herbicides were not used. Notes: See Test 2199 for performance under virus yellows conditions and Test 5399 for performance under

PERFORMANCE OF LINES UNDER RHIZOMANIA, SALINAS, CA., 1999 TEST 5399.

2.3 86.5 85.5 84.5 84.6 87.1 85.0 83.8 85.3 85.8 85.6 84.4 83.8 84.8 84.4 85.4 1.9 87.9 87.1 Harvested: October 28, 1999 RJAP April 29, 1999 2.4** 64.6 18.8 14.6 13.0 18.9 15.2 9.7 11.3 10.6 7.0 23.9 21.7 19.3 Root 8.5 8.9 20.8 Rot 12.7 oo | 5.8** Planted: 8.8 15.5 178.1 Seets/ 1001 190 204 173 182 186 169 181 174 181 182 141 169 183 185 182 162 No No 19.92** Sucrose 14.16 3.20 16.05 16.19 0.52 16.00 15.93 16.84 15.94 15.99 17.61 16.33 15.85 16.65 16.44 16.60 16.24 PERFORMANCE OF LINES UNDER RHIZOMANIA, SALINAS, CA., 1999 11.27** 23.37 28.19 27.65 17.58 22.43 22.56 23.32 25.05 24.79 2.74 11.19 Tons 28.67 23.52 29.85 29.14 21.72 25.78 22.74 25.01 Beets Acre Yield 48 entries x 8 reps., RCB(e), 3 sub-sets of 16 x 8, RCB(e) 15.3** 926.6 11.9 3127.2 Sugar 9868 7258 7685 6686 7560 4985 7467 7443 7150 7532 8413 3984 3029 8241 10140 10263 Lbs Inc. R776, R781, R781-43, ... RZM Y775, Y773, Y772, Y767 RZM R776, R781, R781-43, ... RZM (C80 x EL-smooth root) RZM-8S R576-89-5NB Description Betaseed, 1-19-99 Betaseed, 1-10-99 (690) RZM Y772, (C72) susc. check RZM-8S Y672 RZM Y769, 1-row plots, 22 ft. long RZM R778% RZM Y775 RZM R780 RZM Y768 RZM Y771 MM,O.P. lines 98-EL-04/02 R876-89-5NB R878% (C78) (C82) (C82) TEST 5399. Variety (080) (690) Y872 (C72) Y875 (Sp) LSD (.05) C.V. (%) 5399-1: F value B4776R B4430R US H11 Y872B R881 X869 R882 R880 **X868 Y875** Y871 Mean

85.0 2.0 2.4 2.5**

2.4**

3.6**

3.17

7.31**

10.1**

12.6

63.8

16.4 10.3

> 14.9 8.5

0.51

2.95

9.676

LSD (.05)

Mean

C.V. (%)

F value

24.28

7897.5

16.23

176.9

48 entries x 8 reps., RCB(e). ANOVA to compare means across sets of entries.

TEST 5399. PERFORMANCE OF LINES UNDER RHIZOMANIA, SALINAS, CA., 1999

(cont.)

		Acre Yield	ield		Beets/	Root	
Variety	Description	Sugar	Beets	Sucrose	100'	Rot	RJAP
		Ibs	Tons	ઝ∘	No.	&	₩
5399-2: MM 1	MM lines with Bvm germplasm						
ŀ	Spreckels, 2-8-99	7632	ω.	•	181	6.5	85.3
B4035R	Betaseed, 7-10-97	8518	25.16	16.92	187	11.8	86.7
R827 (C27)	RZM R727A, B	6781	1.0	16.09	173	15.1	86.2
7747	Inc. 5747 (A,aa)	5782	19.64	14.71	177	17.3	86.8
R824	RZM R724, R725 (C79-2/3, WB41, 42)	6123	19.24	15.93	172	19.2	83.3
R835		6545	20.38	16.11	186	10.1	85.1
R879	RZM R779 (C79-1, Rz)	5399	18.25	14.81	163	23.7	84.6
R836	RZM R736, R746 (C79-8, R22)	7152	6.	14.88	181	14.0	81.1
R853	RZM-ER-%S R653, (BC ₄)	6736	21.26	15.81	175	17.6	85.2
R854	RZM R754, (BC ₅)	7254	23.11	15.69	178	18.6	85.4
X873	RZM-ER-%S Y673	7786	23.87	16.35	189	14.9	84.1
X873B	RZM Y773	7134	22.29	16.00	174	13.2	84.6
R840	RZM R740 (C79-#s)	8763	27.07	•	178	11.0	83.8
P811	RZM-PMR 6203-6208-#(C)	7681	24.16	15.88	186	32.5	83.3
X866	RZM Y766	9013	27.45	16.41	181	9.6	84.1
X867	RZM Y767, (C67)	9233	27.47	16.80	182	15.1	85.2
Mean		7345.7	22.95	15.96	179.0	15.7	84.7
LSD (.05)		950.3	6.	•	14.2	11.5	2.3
C.V. (%)		13.1	13.17	3.61	8.0	74.0	2.8
F value		11.2**	7.36**	11.00**	1.8NS	2.3**	2.9**

PERFORMANCE OF LINES UNDER RHIZOMANIA, SALINAS, CA., 1999 TEST 5399.

(cont.)

		H G	Yield		Beets/	Root	
Variety	Description	Sugar	Beets	Sucrose	1001	Rot	RJAP
		Ips	Tons	o⁄e	No.	o,	op
5399-3: MM,	S ^f , Aa populations						
Y869H31	12.	8538	26.21	16.27	173	18.0	84.2
Rifle	Spreckels, 2-8-99	8294	23.34	17.80	170	27.5	85.3
8931	RZM 7931, 6915, 6925(C) aa x A	9968	27.26	16.45	170	15.3	•
Z831	RZM Z731, Z730, Z725(C)aa x A	8820	26.33	16.74	166	18.4	85.6
8924	RZM 7924, aa x A	8841	26.83	16.52	184	14.9	85.1
8926 (Sp)	7931aa x RZM 7926	9223	28.59	16.14	176	17.4	84.3
	RZM 7926aa x A	8440	9	16.01	187	19.6	85.1
8932M	7932ст, 7201-7215Маа х А	7101	21.77	16.34	176	28.6	84.7
P812	RZM-PMR 6211-# - 6217-#(C)	7822	24.40	16.05	179	19.1	83.9
CR811	RZM CR711, (CR09/10)	8222	25.60	16.02	181	9.4	85.1
CR812	RZM CR712	7803	23.92	16.29	171	15.1	84.8
CR813	RZM CR713	8200	25.94	15.82	159	9.4	84.4
N730	Inc. N629, N630 (galls)	7780	24.47	15.91	168	22.7	84.8
8935	RZM R776-89-5H13	7205	21.64	16.65	179	17.5	85.4
8936	RZM R776-89-5H31	8184	24.30	16.84	184	14.7	84.0
8939	RZM Y769H31	8074	24.47	16.50	155	24.8	86.2
Mean		8219.5	25.09	16.40		18.3	84.9
LSD (.05)		882.9	2.66	0.42	15.5	9.5	1.8
C.V. (%)		10.9	10.71	2.57	0.6	50.8	2.2
F value		3.6**	4.03**	10.34**	2.6**	2.8**	0.9NS

Rotted roots weighed at harvest but not included in sugar sample. Root rot caused considerable variability in plot Notes: Root rot due to Sclerotium rolfsii. Roots with obvious rot counted before harvest. weights.

TEST 2999. EVALUATION OF MONOGERM LINES & POPULATIONS, SALINAS, CA., 1999

24 entries x 4 1-row plots, 2	4 reps,sequential 21 ft. long			Planted: 1 Harvested:	March 24, 1999 September 20	99 20, 1999
Varioty	Description	Acre	Yield Reets	esonone	Beets/	RIAP
		Ibs	Tons	₩	No	%
Checks		200	c	c u	7 1	0 2
Kille B4776R	Spreckers, 9-10-90 Betaseed 4776.7653, 3-27-98	14582	41.90	17.44	164	
Monogerm popul	ations					
8810M RZ	RZM 7810NB, (C790 x C890-#)	10204	6.	.5	158	82.7
8833	RZM, T-O 7833-#, 7834-# (A,aa)	8626	27.90	15.45	2	82.0
8835	7835,aa x A	12005	4.	5.6	151	82.5
8836	T-0 7836-#, 7837-# (A,aa)	10193		6.0	2	0
8838	A	11254	35.70	15.75	155	82.5
8848M	RZM 7848, (C790 x C890-#)	10799	34.00	5.9	7	82.8
7869NB	NB-RZM 5869	11490	35.10	6.3	163	82.7
8890	RZM 7890(A,aa), (C890-1Rz)	10602	9	5.7	158	81.4
8932M	7932CT,aa x A	12032	37.10	16.27	152	2
8932н69	6869mmaa x 7932CT,	12984	7	5.7	165	82.2
Monogerm lines	F. hvbrids					
1		11532	36.30	15.95	173	81.8
8829-3H50	ω,	12910	39.40	16.40	167	81.1
8831-3H50	'n	12851	40.30	5.9	163	•
8831-4MHO	‡, (C831-	13436	.5	5.8	154	80.5
8833-5но	C790-15CMS \times 5833-5, (C833-5)	12326	36.00	17.13	161	82.3
8833-12H50	C790-15CMS x 5833-12, (C833-12)	12907	9.7	6.2	2	4.

1999 TEST 2999. EVALUATION OF MONOGERM LINES & POPULATIONS, SALINAS, CA.,

(cont.)

Varieto	Description	Acre Yield Sugar Beet	Vield Beets	Sucrose	Beets/	ሚያ ነ
		Ibs	Tons	~	No.	opo
CMS - monoderm nonlations	populations					
8833H50	C790-15CMS x RZM, T-0 7833-#	11805	36.00	16.41	159	83.3
8835H50	C790-15CMS x 7835	11592	36.30	15.95	158	82.4
8838H50	C790-15CMS x 7838	12402	38.70	16.02	168	82.7
8836MHO	7838H10 x T-O 7836-#, 7837-#	12976	41.50	15.60	158	82.8
8848HO	7848H88 x RZM 7848	11342	34.70	16.36	164	84.8
0н6988	7869HO x RZM 7869-#(C)	11124	34.92	15.91	167	83.1
Mean		11889.1	36.90	16.11	160.7	82.7
LSD (.05)		1596.3	4.80	0.99	15.1	2.5
C.V. (%)		9.5	9.22	4.34	6.7	2.2
F value		5.2**	4.38**	2.03*	1.3NS	2.2**

TEST 4999. RHIZOMANIA EVALUATION OF MONOGERM POPULATIONS, SALINAS, CA., 1999

Planted: April 29, 1999 Harvested: November 1, 1999 24 entries x 8 reps., RCB(e) 1-row plots, 22 ft. long

		Acre	Acre Yield		Beets/	Root	
Variety	Description	Sugar	Beets	Sucrose	100'	Rot	RJAP
		Irbs	Tons	₩	No.	o⁄e	%
Checks		((,	Ç		
Rifle	•	068/	67.77	٠,	181	•	84.3
B4776R	Betaseed, 1-19-99	9886	7 . 4	18.01	184	7.6	87.1
Monogram of	a De Donilations						
3	8700-0-0-0-0-0-0-0-0-0-0-0-0-0-0-0-0-0-0	9065	17 68	14 90	148	7 8 1	86.1
	0.50 51(c) aa A A)	•	•	۲		
8808 (S ₁ C)	RZM-% 68088 (Comp 1-10)	5928	٦.	16.38	162	17.0	84.3
8810M	RZM 7810NB, (C790 x C890-#s)	5745	æ	٥.	173	10.8	84.4
8848M	RZM 7848, (C790 x C890-#s)	6168	19.06	16.19	180	21.1	84.3
8890	RZM 7890 (A,aa), (C890-1Rz)	6373	19.23	16.58	166	20.3	84.9
8833	RZM, T-O 7833-#, 7834-# (A, aa)	5597	17.54	15.98	171	4.4	83.7
8836	T-O 7836-#,7837-# (A,aa)	5450	17.46	15.54	169	14.6	83.8
7869NB	NB-RZM 5869	7167	21.20	σ.	188	11.2	
8835	7835,aa x A	6515		16.50	188	14.1	84.5
8838	7838,aa x A	6460	19.94	6.2	172	14.1	85.1
8932M	7932CTaa x A	6091	٦.	16.38	175	20.5	4.
8932H38	7838mmaa x 7932CT	6845	1.0	16.29	173	21.7	85.3
8932H69	6869mmaa x 7932CT	6882	÷.	•	180	17.0	4.
он6988	7869HO x RZM 7869-#(C)	6625	ω.	9.9	180	17.0	85.4
7818/2M	RZM 6818M (A,aa)	4858	5.5	5.7	177	9	4.
8835H50	C790-15CMS x 7835	6095	18.74	16.27	168	16.9	85.5
8838H50	C790-15CMS x 7838	5958	18.53	ი.	180	24.8	85.3

RHIZOMANIA EVALUATION OF MONOGERM POPULATIONS, SALINAS, CA., 1999 TEST 4999.

(cont.)

		Acre Yield	ield		Beets/	Root	
Variety	Description	Sugar	Beets	Sucrose	100'	Rot	RJAP
		Ibs	Tons	ø•	No.	એ ં	o 0
CMSs of Monogerm Lines	yerm Lines						
8829-3H50	$C790-15CMS \times 5829-3 (C829-3)$	5785	17.11	16.94	178	20.7	83.6
8831-3H50	$C790-15CMS \times 5831-3$ (C831-3)	7823	23.22	16.86	171	27.8	85.4
8831-4MHO	$6831-4MHO \times 7831-4-#s$ (C831-4)	8597	25.48	16.88	164	14.4	83.6
8833-5H50	$C790-15CMS \times 5833-5 (C833-5)$	8585	24.23	17.73	175	23.3	84.4
8833-12H50	C790-15CMS x 5833-12 (C833-12)	7889	23.60	16.71	173	21.5	86.2
Mean		6688.3	20.19	16.48	174.0	17.8	84.8
LSD (.05)		1033.9	2.97	0.61	14.9	11.7	2.0
C.V. (%)		15.7	14.89	3.75	8.7	66.5	2.4
F value		10.8**	7.58**	10.06**	2.7**	1.9*	1.5NS

Notes: Root rot due to Sclerotium rolfsii. Roots with obvious rot counted before harvest. Rotted roots weighed at harvest but not included in sugar sample. Root rot caused considerable variability in plot weights.

TEST 2299. PERFORMANCE OF HYBRIDS UNDER VIRUS YELLOWS INFECTION, 1999

48 entries x 8 reps, RCB(E)	Planted: March 22, 1999
1-row plots, 21 ft. long	Harvested: October 4-5, 1999
	Inoc. BYV/BChV/BWYV: June 22, 1999

			Acre	re Yield			Beets/					
	Variety	Description	Sugar	Loss	Beets	Sucrose	100'	RJAP	Vi	Virus Ye	Yellows	
			Irbs	≫	Tons	. o∤o	No.	₩	07/21	80/80	08/24	Mean
	2299-1: Exper	Experimental hybrids										
	KW6770	Susc.ck, 6770.5193,1-10-97	9951	ე		16.95	168	85.4	4.1	4.8	7.3	5.6
	Rifle	Spreckels, 2-8-99	11253	17.2	34.76		159	2	•	5.0	7.1	5.5
	R876-89-5NBH50	C790-15CMS x RZM-%S R576-89-5NB	16	9	5.9	16.21	164	83.3	3.4	4.9	5.3	•
	R876-89-5H50	C790-15CMS x RZM-%S R576-89-5	11767	4	36.40	6.1	163	•	•	4.6	4.4	4.3
	R882H50	C790-15CMS * R781, R776	11972	14.3	8.4	15.60	9	•	•	4.8	5.0	4.5
	X869H50	C790-15CMS x Y769	148	4.	35.75	0.	156	84.7	3.8	4.6	4.9	4.4
	X868H50	C790-15CMS x RZM Y768	11411	•	5.4		9	ω.	•	5.0	•	4.3
	R878H50 (Iso)	C790-15CMS x RZM R778%	11438		9.	6.0	9	84.0	•	•	5.5	4.7
Δ.	VRKGH50	790-150MS * RZM V766	11047		٧	•	170		۸ ب	ر د		α
49	VR67H50	: >	י מ	- 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1	37.67		1 1 2) () C	•	, r	•
	V871450	; >	16	•	. o	• a	171) (י ר	•	•	•
	0011101	V Part L	14004	5 ι		י טור	7/7	·	0.0	1 , 1	•	•
	Y872H50	C790-15CMS x RZM-%S Y672	Н.	ω.	œ	5. 5.	164	•	•	5. 3	5.4	4.8
	Y872BH50	C790-15CMS x RZM Y772	13	7.7	7.5	0.	9	84.1	4.3	4.5	6.0	5.0
	Y875H50 (Iso)	C790-15CMS x RZM Y775	11735	12.5	•	16.11	167	4.	3.8	4.6	4.9	4.4
	Y873BH50	C790-15CMS x RZM Y773	03	21.7	3.7	5.4	9	82.6	4.1	4.3	5.9	4.8
	R854H50	C790-15CMS x RZM R754	16	•	6.8	5.8	9	m		•	•	4.9
	Mean		11503.8	ļ. I	35.97	16.01	165.1	83.5	3.8	4.7	5.5	4.7
	LSD (.05)		1003.4	ļ. !	ω.	0.57	9.3	•	•	•	9.0	0.4
	C.V. (%)		8.8	. -	8.08	3.62	5.7	2.5	17.1	13.2	10.9	7.7
	F value		* 0.	l	5.17**	2.85**	٠.	NS2.0*	1.7NS	1.3**	15.3**	8.7**
	TEST 2299. PER	REORMANCE OF HYBRIDS UNDER VIRUS FEDS. RCB(E). ANOVA across test	YELLOWS INFE	INFECTION	1, 1999							
			1153	١.	സ	15.95	161.5	83.6	9. 8.	4.7		4.7
			011.	1,	•	. 5	10.	1.8		•	9.0	•
	C.V. (%) F value		ص *	 *	8.33	3.27		•	16.3	13.6 1 2NS	11.	8.2
			•	•	•			•	•	•	,	•

TEST 2299. PERFORMANCE OF HYBRIDS UNDER VIRUS YELLOWS INFECTION, 1999

		Acı	Acre Yield	r		Beets/					
Variety	Description	Sugar	Loss	Beets	Sucrose	1001	RJAP	- 1	0	10w	
		Ips	oke	Tons	≫	<u> </u>	o/P	07/21	08/03	08/24	Mean
2299-2: Hybr	Hybrids with populations										
B4776R	Betaseed, 1-19-99	245	•	6.7	6.9	9	ъ.	3.8	4.8	7.0	
SS-432R	Spreckels, 2-8-99	œ	14.2	33.65	16.13	160	83.4	4.5	4.9	6.8	5.4
8913-70H50	C790-15CMS x RZM-ER-% 6913-70	112	•	4.7	5.9	9	e.	3.3	4.8	4.6	
R882H38	7838mmaa x R781, R776	122	ij.	5.8	5.6	4	4.	•	4.8	•	4.6
8931H50	C790-15CMS x RZM 7931	165	•	37.50	.5	172	δ.		4.4		4.5
8924H50	C790-15CMS x RZM 7924	183	•	7.3	5.8	167	4.				4.7
8932H50	C790-15CMS x 7932CT,		10.7	36.65	15.86	170	83.1	3.9	5.1	5.6	4.9
Z831H50	C790-15CMS x RZM Z730,Z731	112	ij.	4.9	5.9	165	ë.	•	4.8	•	4.8
8926H50 (Sp)	C790-15CMS x RZM 7926	11602	12.7	6.3	5.9	170	ω.	4.3	5.0	5.3	4.8
8935H50 (Iso)	x RZM	160	ά.	5.7	6.2	9	7		4.4	•	4.3
8936H50	C790-15CMS x RZM R776-89-5H31	12740	10.4	39.50	16.13	163	83.3	3.1	•	5.1	•
8937H50	C790-15CMS x RZM R776-89-5H11	46	·	8.3	6.2	9	4.	•	4.3	•	4.4
8938H50	C790-15CMS x RZM Z731H11		0	8.5	15.84	168	8	3.8	4.4	•	4.6
8939H50	x RZM	11611	15.7	37.35	15.55	163	84.0	4.3	4.5	5.1	4.6
CR812H50	C790-15CMS x RZM CR712	190	4.	7.1	15.99	170	4.	3.9	4.5	6.3	4.9
CR813H50	C790-15CMS x RZM CR713	12147	•	9.0	5.5	167	m.	•	4.6	•	4.7
Mean		11762.3	ļ. I	36.83	15.97	•	83.6	•	•	•	•
LSD (.05)		•	ļ. 1	. 1	0.48	10.7	1.7	9.0	0.7	9.0	0.4
C.V. (%)		9.1	ļ. I	8.69	3.02	。	2.0	•	14.1	11.4	9.8
F value		2.0*	 - 	2.02*	4.03**	2.2	**1.6NS	S 2.9**	1.2NS	8.3**	5.8**

PERFORMANCE OF HYBRIDS UNDER VIRUS YELLOWS INFECTION, 1999 TEST 2299.

		Ac	Acre Yield	7		Beets/					
Variety	Description	Sugar	Loss	Beets	Sucrose	1001	RJAP	Ν	Virus Ye	Yellows	
		Tps	%	Tons	₩	No	% ∣	07/21	08/03	08/24	Mean
2299-3: Top	Topcross hybrids										
B4035R	Betaseed, 7-10-97	10762	9	34.00	5.8	165	ω.	4.4	4.4		•
B4419R		11993	9	6.8	16.26	9	85.0	3.5	4.6	6.8	5.2
8931H38	7838mmaa x RZM 7931	11520		36.20	5.9	156	4.	•	4.6	5.4	•
8935H38	7838mmaa x R776-89-5H13	11989	6.1	37.45	16.01	5	84.1	3.6	4.4	4.8	4.3
X869H38	7838mmaa x Y769	11408	10.3	. 7	5.9	Ŋ	83.5	4.1	4.4	5.0	4.4
X869H35	7835aa x Y769	11444	o.	6.3	15.76		2	4.3	4.8		4.9
869Н69	7869aa x Y769	9961	24.5	32.83		157	83.2	•	4.9	5.3	4.7
X869H46	7869-6но × Y769	10941	4.	5.0	15.63	161	83.9	3.6	4.4	•	4.5
Y869H4	C831-3aa x Y769	8	9	2.1	5.6	131	4.	4.1	4.3		4.5
X869H5	C833-5aa x Y769	11736	•	6.8	5.9	Ŋ	2	•	4.3	5.9	•
Y869H12	C833-12aa x Y769	11246		35.22	15.93	135	84.3	4.3	4.6	0.9	5.0
Y869H27	C831-4HO x Y769	11810	14.5	7.1	5.9	Ŋ	84.4	ж. Ж.	4.8	4.6	4.2
Y869H29	C829-3aa x Y769	11309		4.7	16.27		82.7	3.1	4.5	0.9	4.7
Y869H45	C867-1HO x Y769	11309	8.7	35.85		155	83.6	4.3	4.8	5.6	4.9
X869H7	C911-4-7HO x Y769	11785		6.5	٦.		ъ Э	3.5	4.6	4.9	•
R882H27	C831-4HO x R781, R776 (C82)	12057	12.4	8.4	15.70	Ŋ	83.2	3.6	5.1	5.3	4.7
Mean		11333.1	1.	35.71	15.87	153.8	83.7	3.8	4.6	5.5	4.7
LSD (.05)		•	<u> </u> .	2.94	4.	•	1.	0	9.0		0.4
C.V. (%)		8.e 7.e	 *	8.31	2.78	9 v	**1.	14.9	14.2 1.1NS	010	7.8
מבות אינים		•	•)		•	•	•	•		•

Relative % loss values were Notes: Test 2299 was inoculated with virus yellows (BYV-BWYV-BChV) on June 22, 1999. calculated in comparison to noninoculated companion Test 2599 (see Test 2599).

TEST 2099. PERFORMANCE OF S1 TOPCROSS HYBRIDS UNDER VIRUS YELLOWS INFECTION, 1999

Planted: March 22, 1999 24 entries x 4 reps, RCB(E) 1-row plots, 21 ft. long

1-row plots, 21 ft. long	21 ft. long				Harves Inoc.	Harvested: October Inoc. BYV/BWYV/BChV	ted: October 4 BYV/BWYV/BChV:	4, 1999 7: June	9 e 22,	1999
Varietv	Description	Acre	Yield	Sucrose	Beets/ 100'	RJAP		Virus Y	Yellows	***
		sqT	Tons	or 1	No.	₩	08/02	08/16	08/24	Mean
Checks Rifle	Spreckels, 9-98, L1162401	9866	•		170	84.6	5.0		7.8	6.3
B4776R Y869H50	Beta 4776R.7653, 3-27-98 C790-15CMS x Y769	11277 11154	33.50	16.84 16.04	159 158	86.2 83.6	4.8	5.8	7.5	6.0
Topcross to S Y869H69	S ₁ 's from popn-869 7869aa x Y769	10644	33.70	15.79	156	84.0	5.0	5.0	6.3	5.4
Y869H46	69/X × 0H9-698/	10410	32.90	15.79	159	84.4	4.8	5.0	6.3	5.3
х869н69-2	7869-2aa x Y769	10268	3.8	δ.	Ŋ	83.2	4.0	4.5	5.3	4.6
X869H69-4	7869-4aa x Y769	10622	ω.	15.95	146	84.3	5.0	5.3	8.9	5.7
Y869H69-20A	7869-20aa x Y769	9902	32.30	15.34	150	83.9	•	•	6.3	5.5
Y869H69-20B	7869-20Baa x Y769	9676	30.80	15.69	142	84.3	5.0	4.8	5.8	5.2
Ү869H69-24	7869-24aa x Y769	10500	32.60	16.10	158	85.0	4.5	4.5	0.9	5.0
Topcrosses to	to S_1 's from popn-833									
х869н5	5833-5aa (C833-5) x Y769	11727	9		137	2.	•			•
¥869H33-3	7833-3aa x Y769	10614	34.70	15.32	157	83.8	5.0	ა შ	9	5.7
X869H33-10	7833-10aa x Y769	11237	•	9	165	m	4.3	4.3	5.0	4.5
х 869н33-12	7833-12aa x Y769	10466	33.15	15.80	148	84.0	5.3	•	7.3	6.1

(cont.)

		Acre	Yield		Beets/					
Variety	Description	Sugar	Beets	Sucrose	100'	RJAP	Λ	Virus Yellows	Yellow	S
		sqT	Tons	o(r)	No.	e*1	08/02	08/16	08/24	Mean
Topcrosses t Y869H27	to S ₁ 's from popn-831-4 6831-4HO(C831-4CMS) x	x Y769 11137	35.30	15.79	150	83.4	4.0	4.3	5.3	4.5
Y869H27-1	7831-4-1 x Y769	9617	30.70	15.69	129	83.1	4.8	5.3	6.0	5.3
Y869H27-7	7831-4-7aa x Y769	11481	36.40	15.77	161	82.1	4.0	4.8	5.8	4.8
X869H27-8	$7831 - 4 - 8aa \times Y769$	10438	33.30	15.66	\mathbf{c}	•	4.8	5.3	6.5	5.5
Y869H27-10	7831-4-10aa x Y769	11896	37.13	16.04	149	82.3	3.8	4.3	5.8	•
Topcrosses t Y869H34-2	to S ₁ 's from popn-834 7834-2aa x Y769	9184	28.60	16.08	145	83.3		5.0	0.9	5
Y869H34-8	7834-8aa x Y769	0686	31.	ъ.	4	m.	5.3	•	•	•
Topcrosses t Y869H36-3	to S ₁ 's from popn-836 7836-3aa x Y769	9585	30.70	15.64	132	84.0	4.3	5.0	6.5	ۍ
Topcrosses t Y869H79-2	to S ₁ 's from popn-839 7839-2aa x Y769	10419	33.60	15.54	154	84.1	4.8	5.0	6.3	5.3
X869H79-3	7839-3aa x Y769	9839	31.80	15.49	159	84.7	5.8	5.8	7.3	6.3
Me G		10493	7 33 14	15.84	151.7	σ α	7 4	0	ر ب	r m
LSD (.05)		927.0	2	2.49	20.3	H		0.7		0.5
C.V. (%)		9	.3 5.77	1.76	•	1.3	11.5	10.3	7.9	8.9
F value		4	.9**4.93**	7.73**	2.0*		4.6**	4.8**	œ.	9**10.1**

Notes: See Tests B799, 999, 2399, & 5599. These entries tested under virus yellows are an abbreviated list from the above tests.

TEST 2599. PERFORMANCE OF HYBRIDS, SALINAS., CA, 1999

48 entries x 8 reps, RCB(E) 1-row plots, 21 ft. long

Planted: March 24, 1998 Harvested: September 24, 1999

		Acre Yield	eld		Beets/	Root	
Variety	Description	Sugar	Beets	Sucrose	1001	Rot	RJAP
		I.bs	Tons	o%	No.	96	o40
2599-1: Experi	Experimental hybrids					l	I
KW6770	Susc. check, 6770.5193, 1-10-97	13315	36.35	18.36	158	0.0	85.5
Rifle	Spreckels, 2-8-99	13594	38.60	17.60	150	0.4	84.5
R876-89-5NBH50	×	14029	40.70	17.24	158	0.0	4.
R876-89-5H50		13706	41.10	16.67	162	0.0	83.0
R882H50	C790-15CMS x R781, R776, (C82)	13968	43.25	16.14	158	0.0	85.3
X869H50	C790-15CMS x Y769, (C69)	13491	41.35	9		•	5.
X868H50	C790-15CMS x RZM Y768	13173	42.30	15.57	160	•	•
R878H50 (Iso)	C790-15CMS x RZM R778%, (C78)	14199	41.96	16.91	163	0.0	5
X866H50	x RZM 3	14077	41.70	6.8	162	0.3	85.3
X867H50	$C790-15CMS \times RZM Y767$, (C67)	13802	41.35	16.73	158	0.0	4.
X871H50	C790-15CMS x RZM Y771	13823	42.90	Н	163	0.0	85.0
Y872H50	C790-15CMS x RZM-%S Y672, (C72)	14111	42.90	16.45	155	0.0	84.2
Y872BH50	C790-15CMS x RZM Y772	12987	39.30	16.49	162	0.0	84.3
Y875H50 (Iso)	C790-15CMS x RZM Y775	13411	41.30	16.25	159	0.0	
Y873BH50	C790-15CMS x RZM Y773	13275	41.40	0	157	0.0	83.9
R854H50	C790-15CMS x RZM R754	12768	39.55	6.1	155	0.0	
Mean		13608.1	41.00	16.62	158.3	0.1	84.6
LSD (.05)		•	2.49	0.59	10.6	0.5	1.9
C.V. (%)		7.5	6.14	3.58	6.7	671.9	2.2
F value		1.4NS	4.07**	10.51**	23.7NS	0.8NS	1.0NS
2599. ntries x	PERFORMANCE OF HYBRIDS, 1999 8 reps. RCB(E). ANOVA to compare	means across	sets.				
Mean		13493.9	41.08	16.43		0.05	84.5
LSD (.05)		1027.0	2.77	0.58	11.5	0.36	1.7
C.V. (%)		7.7	6.84	3.60	7.5	802.14	2.1
F value		3.1**	3.56**	5.90**	4.8**	0.89NS	1.6**

TEST 2599. PERFORMANCE OF HYBRIDS, SALINAS., CA, 1999

		Acre Yi	Yield		Beets/	Root	
Variety	Description	Sugar	Beets	Sucrose	100'	Rot	RJAP
		Irbs	Tons	o%	No.	æ	o40
2599-2: Hybrid	Hybrids with MM, S ^f , Aa populations						
B4776R	Betaseed, 1–19–99	15006	42.20	17.79	156	0.0	86.7
SS-432R	Spreckels, 2-8-99	12668	38.75	16.34	162	0.0	83.9
8913-70H50		14195	43.55	16.30	164	0.0	83.5
R882H38	7838mmaa x R781, R776	12705	39.85	15.94	148	0.0	•
8931H50	C790-15CMS x RZM 7931	14349	43.90	16.35	156	0.0	84.3
8924H50	×	12849	39.45	16.26	156	0.0	84.8
8932H50	C790-15CMS x 7932CT,	13023	39.80	16.36	166	0.0	83.9
Z831H50	C790-15CMS x RZM Z730, Z731	14103	42.45	16.61	159	0.0	84.1
8926H50 (Sp)	C790-15CMS x RZM 7926	13289	41.45	16.01	156	0.4	84.9
	C790-15CMS x RZM R776-89-5H13	13298	40.80	16.27	156	0.0	85.2
8936H50	x RZM R77	14221	42.40	16.79	161	0.4	83.5
8937H50	C790-15CMS x RZM R776-89-5H11	13880	42.55	16.34	156	0.0	84.4
8938H50	C790-15CMS x RZM Z731H11	13657	42.11	16.21	154	0.0	84.3
8939H50	C790-15CMS x RZM Y769H31	13778	42.75	6.0	161	0.0	84.6
CR812H50	C790-15CMS x RZM CR712	13980	42.75		163	•	83.8
CR813H50	C790-15CMS x RZM CR713	13966	43.50	16.05	165	0.0	84.4
Mean		13685.4	41.77	16.38	158.7	0.05	84.4
LSD (.05)		904.4	2.44	•	8.6	0.38	1.7
C.V. (%)		6.7	5.90	3.78	6.2	801.64	2.1
F value		4.2**	3.29**	3.87**	1.8*	0.91NS	1.6NS

TEST 2599. PERFORMANCE OF HYBRIDS, SALINAS., CA, 1999

			Acre Yi	Yield		Beets/	Root	
Variety	έy	Description	Sugar	Beets	Sucrose	100'	Rot	RJAP
			Ibs	Tons	o(∙	No.	90	90
2599-3:	Topcross hybrids	Ø			i		I	ı
B4035R	Betaseed,	7-10-97	12891	39.10	16.46	166	0.3	83.9
B4419R	Betaseed,	, 1-19-99	14303	42.30	16.91	162	0.0	5
8931H38	7838mmaa	x RZM 7931	13638	42.15	16.17	147	0.0	2
8935H38	7838mmaa	x R776-89-5H13	12763	38.85	16.42	156	0.0	84.5
х869н38	7838mmaa	x Y769	12723	39.55	16.09	148	0.0	85.1
X869H35	7835aa	x Y769	12770	39.45	16.19	159	0.0	
х869н69	7869aa	x Y769	13201	40.45	16.30	157	0.0	
X869H46	0Н9-6987	× Y769	12768	39.60	16.14	153	0.0	84.6
Y869H4	C831-3aa	x Y769	12100	38.27	15.81	111	0.0	84.5
X869H5	C833-5aa	x Y769	14198	42.05	16.90	158	0.0	82.8
Y869H12	C833-12aa	1 x Y769	13745	41.65	16.51	137	0.0	84.8
Y869H27	C831-4HO	x Y769	13809	43.00	16.06	156	0.0	83.8
Y869H29	C829-3aa	× Y769	11925	36.65	16.29	150	0.0	83.0
X869H45	C867-1HO	x Y769	12382	38.00	16.29	147	0.0	4
X869H7	C911-4-7HO x Y769	10 x Y769	14031	43.40	16.14	145	0.0	83.1
R882H27	C831-4HO x R781,	x R781, R776	13766	43.30	15.91	147	0.0	85.2
Mean			13188.3	40.49	16.29	149.9	0.02	84.4
ISD (.05)			1105.0	3.17	0.49	11.0	0.24	1.6
C.V. (%)			8.5	7.92	3.05	7.4	1116.24	1.9
F value			3.6**	3.41**	3.00**	10.8**	0.99NS	2.5**

few root problems were experienced in these trials. Damping-off was not observed and BNYVV and SBCN should Therefore, have been at very low levels. Powdery mildew was controlled as needed as were aphids and other insects accumulate sucrose. Under these conditions, these tests should have been good at measuring the genetic Tests 2099 thru 2999 were grown necessary to use herbicides. The plot area was sprinkler irrigated at least weekly and wilting rarely occurred; i.e. these tests were grown with minimal stress. Under these conditions, the most important It was not factor for sugar yield (other than experimental error) should have been their genetic potential to potential of these materials. Sugar yield of up to 15000 lbs/a for less than 6 months are quite Prior to strawberry, the soil had been fumigated with methyl bromide. Except for mild infestation of black aphids, there was little evidence of insect damage. See Test 2299 for performance under virus yellows conditions. following strawberries. remarkable Notes:

TEST 2699. EVALUATION OF EXPERIMENTAL HYBRIDS, SALINAS, CA., 1999

	8 reps, RCB(E)		Pl	Σ	19	
1-row plots,	21 ft. long		н	Harvested: Sep	September 22-24,	, 1999
		Acre Y	Yield		Beets/	
Variety	Description	Sugar	Beets	Sucrose	1001	RJAP
		I.bs	Tons	%	No.	%
2699-1: Exper	Experimental hybrids with S ₁ pollinators					
SS-432R	Spreckels, 2-8-99	12922	39.40	16.40	157	
Rifle		14374	•	17.29	161	
B4776R	Betaseed 4776R.7653 (3-27-98)	14911	41.35	18.02	156	85.7
8931H50	C790-15CMS x RZM 7931	13288	40.90	16.24	161	•
8925-19H50	C790-15CMS x 6925-19	14773	45.55	16.24	156	•
8913-70H50	C790-15CMS x RZM-ER-%S 6913-70	13953	41.70	16.74	158	•
8911-4-10H50	C790-15CMS x RZM-ER-%S 6911-4-10	14840	•	17.13	159	81.5
8918-12H50	C790-15CMS x RZM-ER-%S 6918-12	14342	44.95	15.96	151	•
8918-21H50	C790-15CMS x RZM 7918-21	14145	44.20	16.01	148	85.1
Z825-6H50	×	15069	•	•	165	83.2
Z825-9H50	$C790-15CMS \times Z625-9$	15044	•	18.21	157	85.1
Z830-11H50	C790-15CMS x Z630-11	14709	45.40	16.23	155	•
R709-1H50	C790-15CMS x CR-RZM R509A-1	14377	42.45		156	83.4
CR812H50	×	13510	•	•	158	•
CR813H50	C790-15CMS x RZM CR713	14182	44.00	16.10	162	84.9
R709-9H50	C790-15CMS x CR-RZM R509A-9	14706	46.45	15.86	165	85.8
Mean		14321.4	43.07	16.65	157.9	84.2
LSD (.05)		1368.7	3.81	•	10.5	1.5
C.V. (%)		9.7	8.94	3.50	6.7	1.8
F value		1.7NS	2.24**	12.15**	1.5NS	4.3**
TEST 2699. E	EVALUATION OF EXPERIMENTAL HYBRIDS, 1999 8 reps, RCB(E). ANOVA to compare means	across sets	of entries			
c		14041.7		16.46	155.7	84.2
LSD (.05)		1321.4	3.61	0.62	•	1.6
C.V. (%)			•	3.84	•	•
F value		2.2**	1.90**	6.17**	1.4NS	2.0**

TEST 2699. EVALUATION OF EXPERIMENTAL HYBRIDS, SALINAS, CA., 1999

(cont.)

Varietv	Description	Acre Y Sugar	Yield Beets	Sucrose	Beets/ 100'	RJAP
		I.bs	Tons	o4∙	No.	o/e
2699-2: S ₁ pol.	S. pollinators from MM, VY, Sf, Aa, Rz popns					
R878H50	C790-15CMS x R778, R778%	13129	40.00	16.41	144	83.8
8930-19H50	C790-15CMS x 6930-19	14744	43.85	16.81	156	84.6
8930-39H50	C790-15CMS x 6930-39	14356	43.95	16.30	160	84.5
8930-102H50	C790-15CMS x 6930-102	14107	42.45	16.64	158	83.1
R882H50	C790-15CMS x R781, R776	14106	44.10	15.96	146	84.7
R876-89-5H50	C790-15CMS x RZM-%S R576-89-5	14662	44.45	16.49	154	84.3
8929-41H50	C790-15CMS x 6929-41	14523	43.60	16.65	159	84.1
8929-72H50	C790-15CMS x 6929-72	14082	43.30	16.25	158	84.8
8929-102H50	C790-15CMS x 6929-102	14039	42.85	16.39	155	83.5
8929-112H50	C790-15CMS x 6929-112	14236	ა.	17.14	159	83.6
8929-114H50	C790-15CMS x 6929-114	14985	45.05	16.63	152	84.3
8929-115H50	C790-15CMS x 6929-115	13970	40.90	17.08	151	84.0
8929-133H50	C790-15CMS x 6929-133	12993	39.30	16.56	152	84.3
8929-153H50	C790-15CMS x 6929-153	13639	41.75	16.34	157	84.6
8929-154H50	C790-15CMS x 6929-154	15489	46.45	16.68	154	83.8
8924H50	C790-15CMS x RZM 7924	14137	42.45	16.65	158	84.3
Mean		14199.9	42.87	16.56	154.5	84.1
LSD (.05)		1204.3	3.18	0.53	10.6	1.5
C.V. (%)		8.6	7.48	3.22	7.0	1.8
F value		2.1*	2.73**	2.49**	1.5NS	0.8NS

1999 EVALUATION OF EXPERIMENTAL HYBRIDS, SALINAS, CA., TEST 2699.

(cont.)

Variety	Description	Acre Y	Yield	Succession	Beets/	0 4
		The	Tons	*		NOA.
2699-3: Lines	& S, pollinators from MM, S ^f , Aa, R22 popns	1		1		۰I
4035R		_ 13763	42.25	16.27	161	84.0
Rizor	Holly HH108, 9-3-97	14558	42.30	17.20	156	84.5
X869H50	C790-15CMS x Y769	14284	43.70	16.33	148	
R835H50	C790-15CMS x RZM R735 (C79-7)	13128	40.45	•	151	84.1
R836H50	C790-15CMS x RZM R736, R746 (C79-8)	13232	41.95	15.79	149	83.8
X873BH50	C790-15CMS x RZM Y773	12854	40.70	15.79	163	84.5
R879H50	C790-15CMS x RZM R779 (C79-1)	12679	41.65	15.23	155	4
X867H50	C790-15CMS x RZM Y767 (C67)	13724	42.60	16.06	155	85.0
Y872H50	x RZM-	13762	43.30	15.90	157	83.8
X875H50	C790-15CMS x RZM Y775,	12596	39.60	15.85	154	85.2
8926H50 (Sp)	C790-15CMS x RZM 7926,	13847	42.70	16.24	151	m.
8926H50 (Iso)	C790-15CMS x RZM 7926	13306	41.95	15.86	159	84.0
8927-29H50	C790-15CMS x 6927-29	14514	42.85	16.91	156	83.9
8927-30H50	× 6927-	13105	40.65	16.15	152	82.1
8927-33H50	× 6927-	13770	41.55	16.55	150	83.8
8927-37H50	C790-15CMS x 6927-37	14542	44.70	16.27	159	85.2
Mean		13603.9	42.06	16.16	154.7	84.1
LSD (.05)		1398.7	3.79	0.67	10.1	1.9
C.V. (%)		10.4	9.11	4.17	9.9	2.3
F value		1.7NS	0.93NS	3.85**	1.5NS	1.3NS

The best S_1 lines were selected, increased in isolation, selfed to produce multigerm, S_1 progeny lines. These S_1 lines were evaluated per se for bolting, tendency, hybrids in tests in Imperial Valley and Salinas, the superior progeny lines will be reselected for further In general, So plants from populations were selected for resistance to rhizomania and/or virus yellows and lines extracted from multigerm, self-fertile, genetic-male-sterile facilitated random-mated populations. and crossed to the tester C790-15CMS to produce testcross hybrids. Based upon the performance of these C790-15CMS was used as a common tester to evaluate the general combining ability of S_1 disease resistance, and components of sugar yield. evaluation, improvement, and recombination. Notes:

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TEST 2899. EVALUATION OF POPULATION HYBRIDS, SALINAS, CA., 1999

24 entries x 8 1-row plots, 21	8 reps, RCB(E) 21 ft. long			Planted: Harveste	Ma d:	rch 24, 1999 September 21,	1999
		Acre	Yield		Beets/	Root	
Variety	Description	Sugar	Beets	Sucrose	1001	Rot	RJAP
		I.bs	Tons	o⊱	No.	% I	∞ 1
<u>Checks</u> Rifle	Spreckels, 9-16-98	13754	38.05	8.0	151	0.7	
B4776R	7	14814	1.7	17.79	162	0.0	84.7
Population hybrids	ids C790-15CMS * R778 R778* (C78)	13532	41 40	16 34	154	c	α α
R878H55		13212	6.	6.1	9	0.0	. 4
R878H58	7838H50 x R778, %	13304	41.40	16.08	140	0.0	83.6
R878H69	7869aa x R778, %	13542	42.25	6.0	152	0.0	84.9
R876-89-5NBH50		13258	4.	16.39	161	0.0	84.4
R882H50	C790-15CMS x R781, R776 (C82)	13330	41.90	5.8	158	0.0	84.2
R882H55	7835H50 x R781, R776	13826	44.45	15.55	155	0.0	84.3
R882H58	7838H50 x R781, R776	13707	43.00	5.9	153	0.4	82.7
X875H55	x RZM	13314	41.35	16.10	155	0.4	84.3
X875H58	7838H50 x RZM Y775,	13291	41.10	6.1	Ŋ	0.0	ë.
8931H50	C790-15CMS x RZM 7931,	14744	45.45	16.21	159	0.0	•
8931H38	7838mmaa x RZM 7931,	13256	41.05	16.16	155	0.0	83.1
8932H38	7838mmaa x RZM 7932CT,	13657	42.80	15.94	151	0.0	82.7
8935H38	7838mmaa x R776-89-5H13	14074	42.55		156	0.0	83.5
8935H50	C790-15CMS x R776-89-5H13	14828		0.	147		
8936H50	C790-15CMS x RZM R776-89-5H31	14338	43.30	6.5	156	0.0	83.2
8937H50	x RZM	13889	43.00	16.13	162	0.0	
8938H5 0	C790-15CMS x RZM Z731H11	14239	43.75	6.2	157	0.0	•

1999 SALINAS, CA., EVALUATION OF POPULATION HYBRIDS, TEST 2899.

(cont.)

		Acre Yield	ield		Beets/	Root	
Variety	Description	Sugar	Beets	Sucrose	100'	Rot	RJAP
		Lbs	Tons	%	No.	₩	o⁄e l
Population hy	Population hybrids (cont.)						
8939H50	C790-15CMS x RZM Y769H31		43.85	15.76	148	0.0	81.4
Z831H50	C790-15CMS x RZM Z730, Z731		44.30	16.29	146	0.0	82.9
Z831H55	7835H50 x RZM Z730, Z731	14067	43.45	16.16	145	0.0	84.3
Z831H58	7838H50 x RZM Z730, Z731		42.20	16.77	154	0.0	83.5
Mean		13848.0	42.38	16.35	153.8	0.1	83.8
LSD (.05)		1152.2	2.94	0.73		0.5	2.2
C.V. (%)		8.5	7.04	4.51	9.6	820.1	2.7
F value		1.6NS	2.23*	4.97**	1.2NS	0.9NS	1.3NS

conventional self-sterile, open-pollinated lines such as C78, C69, etc., or self-fertile, genetic-male-sterile traits of various populations and to determine which populations combine well together. Populations that show good performance may then be chosen as a source of progeny lines, e.g., \mathbf{S}_1 progenies. Various types of intramonogerms in the 800-series, e.g., popn-869. Test 2899's purpose was to determine in general the performance reciprocal recurrent selection) can be done. The major thrust continues to be to develop source populations population improvement (e.g., mass and recurrent selection) and modified interpopulation improvement (e.g., developed. The multigerm populations are generally numbered in the 900-series, e.g. popn-931, and the facilitated, random-mated populations. Both multigerm and monogerm self-fertile populations have been Notes: Much of the breeding program at Salinas involves population improvement. Populations may be with useful combinations of disease resistance and tolerance.

TEST 2799. EVALUATION OF TOPCROSS HYBRIDS, SALINAS, CA., 1999

24 entries x 1-row plots,	8 reps, RCB(e) 21 ft. long			Planted: 1 Harvested:	March 24, 1999 September 21	9 1, 1999
		Acre	Yield	8	Beets/	1
Variety	Description	Sugar	Tons	Sucrose	No.	RJAP *
Checks		0	(1	1	
B4776R Rifla	Betaseed 4776.7653, 3-27-98 Spreckels, 9-16-98	13972	39.45 38.95	17.70	158 156	85.5 84.6
))))) 1	P .
Topcrosses to	released mm lines					
1	C790-15CMS x Y769	13315	42.14	15.75	152	84.3
У869Н 4	5831-3aa (C831-3) x Y769	12235	38.47	Ŋ.	102	83.0
X869H5	5833-5aa (C833-5) x Y769	13584		9	141	83.1
X869H7	1-4-7)	12655	40.75	ഗ	148	83.3
R678H33-5	·5) x R5	14383	•	7.6	134	4.
X869H27	6831-4HO (C831-4) x Y769	13574	42.90	15.79	154	84.7
X869H29	x Y76	12636	•	16.06	145	
Y869H45	7867-1HO (C867-1) x Y769	12726	39.30		146	85.2
х869 H46	7869-6НО х Y769	12556	•	5	148	
Topcrosses to						
X869H17	7817HO (C790-7) x Y769	12565	40.05	15.65	155	85.1
Y869H18	7818HO (C790-8) x Y769	12681	0.2	Ŋ.	156	9
X869H49	7848H88mm x Y769	12312	39.80	15.50	156	84.3
X869H35	7835mmaa x Y769	233	0.6	'n.	144	w.
х869н38	7838mmaa x Y769	12428	ი.	Ŋ.	148	•
X869H55	7835H50 x Y769	12239	9.4	5.4	143	84.5
X869H58	7838H50 x Y769	12196	38.80	9.	158	
X869H69	x Y769	29	7.0	س	161	83.9
хвеэнвв	7890HO (C890-1) x Y769	12140	39.60	. 7	149	

TEST 2799. EVALUATION OF TOPCROSS HYBRIDS, SALINAS, CA., 1999

(cont.)

Variety	Description	Acre Yield Sugar Beet	Tield Beets	Sucrose	Beets/ 100'	RJAP
		I.bs	Tons	o/o	No.	o%
Topcrosses to mm populations (cont. Y869H30M 7932CTMaa x Y769	opulations (cont.) 7932CTMaa x Y769	12733	39.50	16.13	147	83.6
Y869H31	7931aa x Y769	12819	40.55	15.81	145	83.4
C831-4 topcrossed R882H27	6831-4HO (C831-4) x R781,R776	13141	41.80	15.70	150	83.2
X875H27	6831-4HO (C831-4) x Y775	12918	41.45	15.59	147	82.1
Mean		12862.6	40.17	16.00	147.7	83.9
LSD (.05)		1253.2	3.06	1.39	14.8	2.1
C.V. (%)		6.6	7.73	4.94	10.2	2.5
F value		1.8*	1.03NS	6.20**	4.8**	1.8*

TEST 2399. HYBRID PERFORMANCE OF MONOGERM S1 PROGENY LINES, 1999

72 entries x 4 1-row plots, 21	reps., RCB ft. long			Planted: N Harvested:	March 24, 19 September	1999 r 27, 1999
**************************************	Description	Acre	Yield	Sucrord	Beets/	R.TAP
		I.bs	Tons	& I	No.	o/o
<u>Checks</u> Rifle	Spreckels, 9-98, L1162401	36	38.40	4.	155	84.4
B4776R	Beta 4776R.7653 (3-27-98)	Ŋ	42.90	7.	169	85.2
X869H50	C790-15CMS x Y769	78	6.	4.	173	84.8
X869H46		454	43.40	6.8	158	•
S, lines from p	popn-833					
	7835aa xY769	368	1.4	6.5	161	83.2
X869H5	5833-5aa (C833-5) x Y769	13336	38.80	17.25	161	82.4
X869H33-1	7833-1aa x Y769	150	5.6	6.1	164	83.1
X869H33-3	7833-3aa x Y769	275	8.0	6.7	156	m.
х869н33-1 0	x Y7	14527	43.53	16.69	155	83.9
X869H33-11	7833-11aa x Y769	14224	4.0	6.1	163	ω.
х869н33-12	x Y7	12707	39.80	15.94	145	83.6
X869H12	5833-12aa (C833-12) x Y769	11976	6.8	6.2	93	4
S ₁ lines from p	from popn-834					
Y869H29	5829-3aa (C829-3) x Y769	12541	38.53	6.2	145	æ.
Y869H34-1	×	α	8.5	6.4	156	ж
Y869H34-2	7834-2aa x Y769	8	38.40	16.65	158	82.9
Y869H34-3	7834-3aa × Y769	339	40.10	6.7	149	Η.
Y869H34-5	×	13489	. 7	5.7	Ŋ	84.0
Ү869н34-8	7834-8aa x Y769	13107	40.80	16.09	156	84.7
Y869H28-9	x Y76	13242	2	15.48	159	83.8
X869H28-10	7828-10aa x Y769	13665	42.80	16.01	168	84.7

TEST 2399. HYBRID PERFORMANCE OF MONOGERM S1 PROGENY LINES, 1999

Variety	Description	Acre	Yield Beets	Sucrose	Beets/ 100'	RJAP
		Lbs	Tons	o⊱	No.	o% I
S, lines from popn-869	698					
х869Н69	7869aa x Y769	13219	•	6.0	144	
1 -69H69X	1aa x Y76	13448	41.50	6.2	146	82.8
Т869H69- 2	7869- 2aa x Y769	13923	43.78	15.90	154	84.0
¥869H69- 4	4aa x Y76	12530	37.50	6.7	148	82.3
Х869Н69- 5		12721	0	. 7	131	84.3
9 -69Н698Х	7869- 6aa x Y769	12834	•	7.	157	٠
7 -69Н698Т	x Y7	13278	6	16.81	152	•
¥869H69-13	7869-13aa × Y769	14662	•	4.	150	84.2
Y869H69-19	7869-19aa x Y769	13928	4.	6.0	156	•
Ү869H69-2 0	7869-20aa x Y769	13688	41.40	16.56	162	84.0
Y869H69-20B		13485	1.9	6.1	152	•
Ү869H69-24	7869-24aa x Y769	12780	•	9.9	164	•
S ₁ lines from popn-836	836					
Х 869Н38	7838aa x Y769	12862	8.0	15.71	4	ω.
Х869Н36- З	× X76	12829	9.3	6.2	133	α.
х869H36-11	x X76	13477	41.20	16.33	m	
Y869H36-14	7836-14aa x Y769	12558	ä	6.4	133	83.8
S ₁ lines from popn-837	837					
	37-1aa x Y76	13222	40.56	16.31	120	83.2
X869H77-1B	7837-1Baa x Y769	13012	40.00	•	140	
X869H77-2	7837-2aa x Y769	13599	41.80	16.25	156	81.2
X869H77-3	7837-3aa x Y767	13713	42.40	٦.	159	83.1
Y869H77-4	7837-4aa x Y769	12143	37.90	16.01	144	83.7

TEST 2399. HYBRID PERFORMANCE OF MONOGERM S1 PROGENY LINES, 1999

Variety	Description	Acre	Yield Beets	Sucrose	Beets/ 100'	RJAP
		sqT	Tons	∞	No.	₩
S ₁ lines from popn-839	n-839					
Y869H79-1	7839-1aa x Y769	12545	•		168	m.
X869H79-2	x Y7	12537	•		145	83.6
X869H79-3	7839-3aa x Y769	13459	41.10	16.38	162	4
X869H79-4	× Y7	11858	36.00	16.47	163	83.4
X869H79-5	7839-5aa x Y769	13819	42.30	16.31	151	83.4
Y869H79-5B	7839-5Baa x Y769	13888	41.70	16.65	145	83.7
X869H79-6	7839-6aa x Y769	13242	40.40	4.	157	84.9
X869H79-10	7	13886	44.00	15.80	154	
S, lines from pop	popn-831-4					
	5831-3aa (C831-3) x Y769	13614	41.55	•	118	
Y869H27-1	7831-4-1aa x Y769	13113	39.98	16.38	123	
Y869H27-2	×	13739	•	•	129	82.1
Y869H27-7	7831-4-7aa x Y769	14040	42.90	16.39	161	81.8
Y869H27-8	7831-4-8aa x Y769	13328		۲.	138	83.4
Y869H27-9	×	12640	•	16.51	113	81.1
Y869H27-10	$7831-4-10aa \times Y769$	13953	43.10	6.2	140	81.8
S ₁ lines from pop	908-udod					
Y869H9-1	7808-1aa x Y769	13826	42.80	16.14	148	84.2
Ү869Н9-2	7808-2aa x Y769	13911	41.70	16.67	143	83.6
Х869Н9-3	7808-3aa x Y769	13697	7.	æ	132	82.2
Y869H9-4	7808-4aa x Y769	12914	40.20	16.09	154	84.1
X869H9-7	7808-7aa x Y769	13890	43.97	15.81	137	83.0
X869H9-8	× X76	13002	40.78	15.90	125	84.4
X869H9-9	7808-9aa x Y769	12542	۲.	16.46	137	84.3
Ү869 H9-12	7808-12aa x Y769	12633	41.60	15.21	145	83.1

1999 TEST 2399. HYBRID PERFORMANCE OF MONOGERM S1 PROGENY LINES,

(cont.)

		Acre Yield	/ield		Beets/	
Variety	Description	Sugar	Beets	Sucrose	1001	RJAP
		sqT	Tons	· %	No.	₩
S ₁ lines from popn-808 (cont.)	n-808 (cont.)					
<u> Ү</u> 869Н9-13	× Y7	13270	41.50	16.02	146	83.4
Х869H9-16	7808-16aa x Y769	12132	38.50	15.76	125	85.1
S, lines from popn-818	n-818					
X869H15-1B	6818-1Baa x Y769	13119	39.50	16.66	158	83.0
Y869H15-2B	6818-2Baa x Y769	13648	42.20	16.21	145	82.3
X869H15-1	6818-1aa x Y769	13361	40.78	16.41	144	83.0
X869H15-2	6818-2aa x Y769	12745	39.90	15.96	157	82.4
X869H15-6	6818-6aa x Y769	13384	42.00	15.96	163	80.3
X869H15-21	6818-21aa x Y769	12687	39.70	15.97	134	83.4
Mean		13279.1	40.78	16.29	147.8	83.3
LSD (.05)		1899.2	5.51	0.71	15.9	2.3
C.V. (%)		10.3	9.70	3.13	7.7	2.0
F value		1.1NS	s 1.11NS	2.89**	6.6**	1.8**

tester. Selected S1 progeny were topcrossed to C69 using genetic male steriles to produce topcross hybrids for evaluating early generation general combining ability for components of sugar yield. Test 2399 serves as a screening trial to determine if any of these S1 lines are worth further breeding effort. Also, these Notes: Monogerm, self-fertile, genetic-male-sterile facilitated random-mated populations that segregate for resistance to rhizomania, O-type, etc. have been developed. From these, rhizomania resistant (Rz_) aplants were selected, selfed to produce S1 progeny lines, and crossed to an annual, male-sterile, type-O hybrids are used to evaluate the potential value of these source populations for future breeding efforts and population improvement.

PERFORMANCE OF EXPERIMENTAL HYBRIDS, SALINAS, CA., 1999 TEST 5499.

48 entries x 8 1-row plots, 2	8 reps., RCB(e) 22 ft. long				Planted: Apr Harvested: No	April 29, 1999 : November 2, 1	1999
		Acre Yi	eld		Beets/	Root	
Variety	Description	Sugar	Beets	Sucrose	1001	Rot	RJAP
		Tps	Tons	∞ 1	No.	æ1	%
5499-1: Testo	Testcross hybrids	7303	16 10	16 76	16/		0
NW6/ / O		7 7 8 9	· α	17.93	F 24		0 0 0 0
RATE-80-5NBH50	JPIECKEIS, Z 0 33 0 C790-15CMS * RZM-%S R576-89-5NB	7785	22.23	17.50	178		85.7 2
US H11	susceptible check	4188	14.52	14.27	153	36.3	85.1
R882H50	C790-15CMS x R781, R776 (C82)	8004	24.51	9	172	30.7	86.1
Y869H50	x Y769 (C69)	6451	9.3	9	173	31.1	9
X868H50	x RZM Y	5355	16.56	•	185	35.0	86.1
R878H50 (Iso)	C790-15CMS x RZM R778% (C78)	6729	19.84	9	174	38.8	7 .
X866H50	C790-15CMS x RZM Y766	7716	22.77	φ.	181	29.5	85.7
X867H50	C790-15CMS x RZM Y767 (C67)	6461	19.52	16.51	178	40.0	ω.
Y871H50	C790-15CMS x RZM Y771	9271	27.71	ø.	182	33.6	85.4
X872H50	C790-15CMS x RZM-%S Y672 (C72)	9222	27.26	16.90	181	33.5	ė.
Y872BH50	×	8674	5	7.	184	w.	
Y875H50 (Iso)	C790-15CMS x RZM Y775	6884	o.	ė.	180	38.2	
X873BH50	C790-15CMS x RZM Y773	6871	21.08	16.26	178	31.6	87.2
R854H50	C790-15CMS x RZM R754	5671	7.	9	169	5.	
Mean		9.6007	21.00	16.59	174.6	33.5	85.8
LSD (.05)		1240.4	3.69	0.53	17.9	12.9	2.2
C.V. (%)		7.	17.76	3.21	10.3		2.5
F value		10.5**	8.64**	16.88**	1.9NS	1.3NS	1.0NS
TEST 5499. PE 48 entries x 8	PERFORMANCE OF EXPERIMENTAL HYBRIDS, 8 reps., RCB(e). ANOVA to compare:	, SALINAS, CA means across	CA., 1999 oss sets of	entries.			
Mean		7612.1	22.70	16.72	176.9		85.9
LSD (.05)		•	3.79	0.59	8.2	13.3	2.2
C.V. (%)		17.6	16.94	5.98	4.7	45.2	2.6
F value		6.2**	5.72**	6.68**	11.0**	2.0**	1.2NS

TEST 5499. PERFORMANCE OF EXPERIMENTAL HYBRIDS, SALINAS, CA., 1999

		Acre Yield	ield		Beets/	Root	
Variety	Description	Sugar	Beets	Sucrose	1001	Rot	RJAP
		I.bs	Tons	1%	No.	o(0	o/0
5499-2: Popula	Population hybrids						
B4776R	Betaseed, 1-19-99	9703	27.40	17.70	190	14.1	87.0
SS-432R	Spreckels, 2-8-99	6962	20.46	17.06	159	24.1	84.6
8913-70H50	x RZM-ER-	7991	23.77	16.77	185	35.5	86.3
R882H38	7838mmaa x R781, R776 (C82)	8279	24.87	16.65	161	26.2	86.8
8931H50	C790-15CMS x RZM 7931	7730	22.61	17.04	181	31.8	85.4
8924H50	C790-15CMS x RZM 7924	6863	20.78	16.54	179	32.3	86.2
8932H50	C790-15CMS x 7932CT,	6106	18.88	16.26	181	31.8	86.1
Z831H50	C790-15CMS x RZM Z730, Z731	7231	21.65	16.64	178	34.9	86.5
8926H50 (Sp)	C790-15CMS x RZM 7926	7351		16.48	180	34.1	85.2
	x RZM R77	8011	3.2		181	Ä	
8936H50	C790-15CMS x RZM R776-89-5H31	8224	23.92	17.23	173	31.3	83.9
8937H50	C790-15CMS × RZM R776-89-5H11	8080	23.67	17.06	188	29.4	86.8
8938H50	C790-15CMS x RZM Z731H11	6446	18.80	17.17	175	49.1	85.8
8939H50	C790-15CMS x RZM Y769H31	7519	22.50	16.45	181	27.1	84.0
CR812H50	C790-15CMS x RZM CR712	6471	ω.	16.74	195	39.9	86.8
CR813H50	C790-15CMS x RZM CR713	8198	24.93	16.45	7		86.0
Mean		7572.8	22.43	16.84	178.9	31.7	85.8
LSD (.05)		1331.0	3.72	0.68	15.5	12.1	2.2
C.V. (%)		17.8	16.76	4.09	8.8		2.5
F value		3.6**	3.24**	2.55**	2.8*	3.0**	1.6NS

PERFORMANCE OF EXPERIMENTAL HYBRIDS, SALINAS, CA., 1999 TEST 5499.

(cont.)

			Acre Yi	Yield		Beets/	Root	
Variety		Description	Sugar	Beets	Sucrose	1001	Rot	RJAP
			Lbs	Tons	% I	No.	%	o, 0
5499-3:	Topcross hybrids	rids						
B4035R	Betaseed,	ed, 7-10-97	9125	7.2	٠.	196	9	86.9
B4419R	Betaseed,	ed, 1-19-99	6511	18.69	17.42	182	23.8	86.8
8931H38	7838mm	7838mmaa x RZM 7931	8347	5.1	6.5	165	т М	5
8935H38	7838mmaa x	aa x R776-89-5H13	7432		9.9	175	27.4	86.3
X869H38	7838mmaa	aa x Y769	8422	5.5	16.48	166	23.7	87.1
X869H35	7835aa	x Y769	7485	22.69	16.46	183	9	85.0
Х869Н69	7869aa	x Y769	7750	23.33	16.69	182	25.6	
Ү869н4 6	0Н9-6981	но ж ⊻769	8209	24.53	16.73	177	5.	86.1
X875H55	7835H5	7835H50 x Y775	8143	4.5	6.6	176		84.9
8931H46	7869-6	7869-6HO x RZM 7931	7771	3.5	16.49	181	22.9	85.4
Y875H27	6831-4	6831-4HO x Y775	9195	27.32	16.81	170	27.3	85.9
X869H27	6831-4HO x	но х у769	8379	5.6	16.34	175	24.7	85.1
B4430R	Betaseed	pe	84		7.8	0	21.8	87.8
Y869H45	7867-1	7867-1HO x Y769	8171	24.88	16.45	173		5
X869H7	6911-4	6911-4-7HO x Y769	36	24.77	6.8	159	23.5	
R882H27	6831-4HO x	HO x R781, R776	8912	26.73	16.64	171		
Mean			8253.5	24.66	16.74	177.2	24.2	86.0
LSD (.05)	•		88.	2.74	.5	15.5	11.1	2.2
C.V. (%)			12.1	11.23	3.37	8.8	9	•
F value			5.1**	5.34**	3.88**	4.2**	0.6NS	1.2NS

Roots with obvious rot counted before harvest. Rotted roots weighed at harvest but not included in sugar sample. Root rot caused considerable variability in plot Root rot due to Sclerotium rolfsii. weights. Notes:

EVALUATION OF TESTCROSS HYBRIDS, SALINAS, CA., 1999 TEST 5899.

Planted: April 29, 1999 Harvested: November 1, 1999

48 entries x 4 reps., RCB 1-row plots, 22 ft. long

		Acre Yield	rield		Beets/	Root	
Variety	Description	Sugar	Beets	Sucrose	1001	Rot	RJAP
		Lbs	Tons	₩]	No.	o40	o40
Experimental	Experimental hybrids with S ₁ pollinators						
US H11	Susc. check	6054	20.23	0.	155	•	ر ا
Rifle	Spreckels, 2-98, L1162401	8501	23.16	18.40	168	25.3	85.9
B4776R	Betaseed 4776R.7653 (3-27-98)	10973	30.07	7	190	•	87.8
8931H50	C790-15CMS x RZM 7931	8323	25.30	٠.4	186		82.7
8925-19H50	C790-15CMS x 6925-19	10653	0	17.70	186	26.7	7.
8913-70H50		9685		17.63	185	o.	•
8911-4-10H50	C790-15CMS x RZM-ER-%S 6911-4-10	10118	28.00	18.08	175	25.7	83.7
8918-12H50	C790-15CMS x RZM-ER-%S 6918-12	9651		17.83	177	0	88.1
8918-21H50	C790-15CMS x RZM 7918-21	7672	2.5	0.	155	7.	9
Z825-6H50	C790-15CMS x Z625-6	8299	3.6	7.5	165	8	7.
Z825-9H50	C790-15CMS x Z625-9	10250	28.33	18.10	183	32.0	83.7
Z830-11H50	C790-15CMS x Z630-11	8023	4.1	6.5	178	2	5
R709-1H50	C790-15CMS x CR-RZM R509A-1	9039	5.8	17.48	170	⊣	4.
CR812H50	C790-15CMS x RZM CR712	7163	20.62		177	37.0	•
CR813H50		8613	5.8	16.65	180	9	85.4
R709-9H50	C790-15CMS x CR-RZM R509A-9	8379	6.2	5.9	178	Ŋ	85.1
S ₁ pollinator	from MM , VY, S ^f , Aa,						
R878H50 (Sp) C	790-15CMS	7906	2.8	ω.	170	5	ė.
8930-19H50	790-15CMS x	8103	3.3	17.35	169		5.
8930-39H50	C790-15CMS x 6930-39	7617	22.15	7.0	186	32.7	84.9
8930-102H50	C790-15CMS x 6930-102	7884	2.1	17.83	191		5.
R882H50	C790-15CMS x R781,R776	8790	9	ശ	176	19.0	85.7
SS-432R	Spreckels	7983	22.64	17.73	172	13.2	

1999 TEST 5899. EVALUATION OF TESTCROSS HYBRIDS, SALINAS, CA.,

Varietv	Description	Acre Yield Sugar Be	Yield Beets	Sucrose	Beets/ 100'	Root	RJAP
•		Ths	Tons	&	No.	%	o,⇔
S1 pollinators from MM	, VY, Sf, Aa, Rz popns	(cont.)					
8929-41H50	4S ×	9948	29.07	7.1	181	11.1	85.0
8929-72H50	C790-15CMS x 6929-72	6260	œ.	16.88	161	39.1	85.4
8929-102H50	C790-15CMS x 6929-102	8914	5.6	ε.	183	9	5
8929-112H50	×	0	9.7	7.5	173	9	9
8929-114H50	×	11174	31.25	7.	191	13.1	9.98
8929-115H50	C790-15CMS x 6929-115	8811	5.1	7.4	187	2	
8929-133H50	C790-15CMS x 6929-133	9376	.2	ζ.	181	0	85.7
92		7712	2.2	7.3	178	4	
8929-154H50	×	9209	26.19	17.58	192	20.1	83.6
8924H50	C790-15CMS x RZM 7924	7335	1.7	6.7	187	വ	9
S. Mollinatore	re from MM of As R22 noons						
4035R	letaseed, 7-10-97	9144	6.0	7.5	σ	14.7	85.8
Rizor	Holly HH108, 9-3-97	9416	26.07	18.05	198	5	85.4
Y869H50	C790-15CMS x Y769	9054	6.8	6.9	7		85.1
R835H50	C790-15CMS x RZM R735 (C79-7)	8663	5.5	6.9	8	8	5
R836H50	C790-15CMS x RZM R736,R746(C79-8)			16.83	182	15.6	85.4
X873BH50	C790-15CMS x RZM Y773	1960	۲.	6.5	œ	ω.	86.7
R879H50	C790-15CMS x RZM R779 (C79-1)	7003	23.19	15.13	185	35.2	86.5
X867H50	x RZM	7366		6.7	œ	5	5.
Y872H50	C790-15CMS x RZM-%S Y672	10374	1.5		176	5.	4.
875H50	C790-15CMS x RZM Y775,	8442	5.2	16.77	ω	т М	86.5
8926H50 (Sp)	C790-15CMS x RZM 7926,	8367	25.48	16.40	192	20.5	84.6
8926H50 (Iso)	$C790-15CMS \times RZM$	8705	6.1	16.65	œ	9	83.4

EVALUATION OF TESTCROSS HYBRIDS, SALINAS, CA., 1999 TEST 5899.

(cont.)

			Acre Yield	ield		Beets/	Root	
Variety	Description	otion	Sugar	Beets	Sucrose	1001	Rot	RJAP
			Ibs	Tons	o/e	No.	oko [o∳• [
S, pollinator	S, pollinators from MM, S ^f , Aa, R22	Aa, R22 popns (cont.)	ıt.)					
8927-29H50	C790-15CMS x 6927-29		8768	24.06	18.23	165	29.0	85.4
8927-30H50	C790-15CMS x 6927-30	927-30	8156	23.82	17.03	178	21.8	85.3
8927-33H50	C790-15CMS x 6927-33	927-33	9042	25.37	17.80	181	26.4	85.3
8927-37H50	$C790-15CMS \times 6927-37$	927-37	8628	25.65	16.92	165	24.4	85.9
Mean			8683.4	25.26	17.16	179.1	24.5	85.4
LSD (.05)			1938.5	5.18	0.77	26.9	19.5	2.5
C.V. (%)			16.0	15.52	3.19	10.7	57.0	2.1
F value			2.7**	2.08**	6.91**	1.1NS	1.4NS	1.6*

Roots with obvious rot counted before harvest. Rotted roots weighed at harvest but not included in sugar sample. Root rot caused considerable variability in plot Notes: Root rot due to Sclerotium rolfsii. weights.

EVALUATION OF TOPCROSS HYBRIDS, SALINAS, CA., 1999 TEST 5699.

24 entries x 1-row plots,	24 entries x 4 reps., RCB 1-row plots, 22 ft. long			<u></u>	Planted: April 29, Harvested: November	ril 29, 199 ovember 3,	99 1999
		Acre	Yie		Beets/	Root	!
Variety	Description	Sugar	Beets	Sucrose	No.	Rot 1%	RJAP
Checks B4776R	Betaseed 4776.7653, 3-27-98	0266	27.69	18.00	187	11.2	4
Rifle	Spreckels, 9-16-98	7116	19.70	œ	187	28.1	86.3
Topcrosses	to released mm lines						
X869H50	C790-15CMS x Y769	7368	2	9	193	26.6	84.8
US H11	susc. ck.	4468	14.67	15.18	175	7.	87.2
Y869H37	4807HO (C306/2CMS) x Y7		21.35	16.05	194	23.2	85.4
X869H7	6911-4-7HO (C911-4-7) x Y7	752	22.47	16.73	170	23.2	84.0
R678H33-5	×	81	23.05	7.6	170	σ	84.6
Y869H27	6831-4HO (C831-4) x Y7	Y769 8424	24.82	17.00	164	0	•
B4430R	Betaseed	9583	•	18.25	185	•	
Y869H45	7867-1HO (C867-1) x Y7	71	21.40	16.70	180	22.3	86.2
X869H46	7869-6HO × Y7	X769 8500	24.92	17.05	178	18.9	86.9
E							
Y869H17	7817HO (C790-7) x Y7	69 7339	22.45	16.33	173	0	4
х869н18		Y769 7601		6.7	187	24.7	86.1
Y869H49	7848H88mm x Y7			16.65	176		
X869H35	×	X769 8300		17.10	178	13.5	
X869H38	7838mmaa × Y7	Y769 8159	23.87	17.10	162	o.	86.0
X869H55	7835H50 × Y7	84		17.27	180		86.2
X869H58	× 0	X769 7565	22.21	17.02	186	19.2	85.9
X869H69	×	80 1		17.10	189	•	86.6
яянкоях	7890HO (C890-1) × Y7		23.24	16.75	180	•	84.5

(cont.)

		Acre Yield	eld		Beets/	Root	
Variety	Description	Sugar	Beets	Sucrose	100'	Rot	RJAP
		I.bs	Tons	o;e	No.	%	%
Topcrosses t	Topcrosses to mm populations (cont.)	i	i d	L T	Ţ L	0	5
X869H30M	7932CTMaa x Y 769	8591	72.01	11.15	121	7.07	0.4.0
Y869H31	7931aa x Y769	8783	26.26	16.73	173	17.9	83.4
C831-4 topcrossed	ossed						
R882H27	6831-4HO (C831-4) x R781,R776	9906	27.21	16.63	161	16.0	85.6
X875H27	6831-4HO (C831-4) x Y775	8167	24.82	16.45	181	14.9	84.2
Mean		7962.8	23.45	16.93	177.6	21.4	85.6
LSD (.05)		1178.0	3.18	0.79	23.5	15.2	2.7
C.V. (%)		10.5	9.61	3.31	9.4	50.3	2.3
F value		4*8.9	5.86**	5.48**	1.7NS	1.9*	1.5NS

Notes: Root rot due to Sclerotium rolfsii. Roots with obvious rot counted before harvest. Rotted roots weighed at harvest but not included in sugar sample. Root rot caused considerable variability in plot weights.

TEST 5599. HYBRID PERFORMANCE OF MONOGERM S1 PROGENY LINES, SALINAS, CA., 1999

72 entries x 1-row plots,	4 reps., RCB 12 ft. long			дд	Planted: April 29, Harvested: November	19 3,	99 1999
		o l	Yield	(Beets/	Root	(-
Variety	Description	Sugar	Beets	Sucrose	.001	Kot	KJAF
		Tps	Tons	ə ∘	 	*°1	≫
Checks			,	,	,		
Rifle	Spreckels, 9-98, L1162401	8093	22.43	18.02	186	35.6	85.9
B4776R	Beta 4776R.7653 (3-27-98)	9847	8.4	7.3	190	0	7
X869H50	C790-15CMS x Y769	6999	8.0	5.7	176	0	4.
х869н46	7869-6HO x Y769	6634	9.7	6.8	175	m.	9
S ₁ lines from	. popn-833						
	7835aa xY769	9	3.3	Η.	167	6.	7
X869H5	5833-5aa (C833-5) x Y769	9227	26.01	17.73	158	15.2	85.5
Y869H33-1	$7833-1aa \times Y769$	6091	8.3	16.58	177	9	9
хв69н33-3	7833-3aa x Y769	8298	9.	7.5	185	2	9
Y869H33-10	7833-10aa x Y769	7001	9.	7.5	189	9	
X869H33-11	7833-11aa x Y769	6528	9.2	0.	183	6	9
Х869Н33-12	×	8374	•	17.13	153	28.9	85.4
US H11	susc. check	5221	7.9	14.50	175	9	9
S ₁ lines from	popn-834						
B4035R	٠.	7415	22.19	16.75	186	21.8	85.5
Y869H34-1	×	6977	0.8	9.9	œ	2	9
Y869H34-2	7834-2aa x Y769	6424	8.5	7.2	Ω	4.	4.
<u> Ү</u> 869H34-3	7834-3aa x Y769	7429	1.4	7.2	9		ë.
Y869H34-5	7834-5aa x Y769	7	20.27	16.55	181	35.5	86.1
Y869H34-8	7834-8aa x Y769	7307	1.4	6.9	190	7.	5.
S_1 lines from	from popn-828						
Y869H28-9	ď	6853	22.26	15.38	185	38.0	86.7
Y869H28-10	7828-10aa x Y769	5366	6.9	5.8	183	ت	7

TEST 5599. HYBRID PERFORMANCE OF MONOGERM S1 PROGENY LINES, SALINAS, CA., 1999

(cont.)

:	:	Acre	Acre Yield	¢	Beets/	Root	1
Variety	Description	Sugar	Tons	Sucrose	No.	Rot I%	RJAP
S, lines from	698-udod						
69н69		7535	7	6.9	198	8	85.2
Y869H69- 1	7869- 1aa x Y769	8820	26.01	17.00	173	35.0	86.7
Т869H69- 2	2aa x	7510	ω.	7.2	182	9	5
¥869H69- 4	7869- 4aa x Y769	6734	٦.	6.7	185	6	5.
х869н69- 5	7869- 5aa x Y769	8037	3.9	6.8	155	23.7	88.1
9 -69Н698х	7869- 6aa x Y769	7498	2.9	6.3	174	11.5	84.7
Х869Н69- 7	7869- 7aa x Y769	7870	22.76	17.30	170	18.2	
х869н69-13	7869-13aa x Y769	8112	4.0	6.9	184	9	85.2
Y869H69-19	7869-19aa x Y769	6647	0.5	6.1	187	22.5	85.6
Ү869H69-2 0	7869-20aa x Y769	7757	2.9	7.0	177	17.6	4
Y869H69-20B	7869-20Baa x Y769	8051	24.06	16.73	162	40.5	85.2
Х869H69-24	7869-24aa x Y769	5447	5.5	7.4	185	28.1	9
S_1 lines from	from popn-836						
х869н38	7838aa x Y769	œ	4.4	6.1	172	ю Ж	δ.
х869Н36- 3	×	7800	23.20	16.80	131	22.8	85.6
х869н36-11	7836-11aa x Y769	8818	5.9	6.9	144	9	7.
Ү869Н36-14	7836-14aa x Y769	8882	5.7	7.2	131	H.	9
S ₁ lines from	popn-837						
X869H77-1	7837-1aa x Y769	8577	5.0	17.13	134	21.2	9
X869H77-1B	7837-1Baa x Y769	7975	3.8	16.75	148	7.	9
X869H77-2	7837-2aa x Y769	7511	22.53	16.67	174	19.8	83.8
X869H77-3	7837-3aa x Y767	0669	9.0	16.90	190	5.	5.
X869H77-4	7837-4aa x Y769	8317	25.89	16.08	153	27.0	84.6

TEST 5599. HYBRID PERFORMANCE OF MONOGERM S1 PROGENY LINES, SALINAS, CA., 1999

Varietv	Description	Acre	Yield Beets	Sucrose	Beets/ 100'	Root Rot	RJAP
		rps	Tons	o,e	No.	%	% I
S ₁ lines from	from popn-839						
Y869H79-1	7839-1aa x Y769	7633	0.	9.9	187	21.4	ო
X869H79-2	7839-2aa x Y769	7211	22.15	16.30	178	35.1	86.9
X869H79-3	7839-3aa x Y769	7777	2.9	6.9	σ	- i	86.5
Y869H79-4	7839-4aa x Y769	თ	21.07	4.	186	Η.	رى
X869H79-5	7839-5aa x Y769	8112	3.8	17.02	9	•	
X869H79-5B	7839-5Baa x Y769	8150	24.15	16.90	166	14.0	86.4
X869H79-6	7839-6aa x Y769	6812	0.5	16.58	7	50.1	86.1
X869H79-10	7839-10aa x Y769	7913	24.69	16.02	153	16.3	86.2
ß							
Y869H27	6831-4HO (C831-4)	വ	വ	9	9	4	21
Y869H27-1	$7831-4-1aa \times Y769$	8481	24.34	17.42	119	21.8	84.8
X869H27-2	7831-4-2aa x Y769	$\overline{}$	3.7	7 .	m	ω.	7.
Y869H27-7	$7831-4-7aa \times Y769$	− œ	8.4	7.4	7	2.	84.4
X869H27-8	7831-4-8aa x Y769	ഥ	7.6	17.20	9	7.	è.
Y869H27-9	$7831 - 4 - 9aa \times Y769$	9426	27.51	œ.	141	18.3	82.6
Y869H27-10	7831-4-10aa x Y769	ഥ	9.8	7.	9	4.	ъ.
S_1 lines from	808-udod						
Y869H9-1	7808-1aa x Y769	39	4.6	7.0	175	27.8	86.7
Y869H9-2	7808-2aa x Y769	90	0.7	7.0	œ	7.	7.
X869H9-3	7808-3aa x Y769	6945	20.83	•	9	•	4.
Y869H9-4	7808-4aa x Y769	8007	3.1	7.2	9	9	7.
7-6H698Y	7808-7aa x Y769	7855	3.6	6.6	Ŋ	6	ъ.
X869H9-8	7808-8aa x Y769	7123	22.05	16.17	164	19.4	87.2
X869H9-9	7808-9aa x Y769	7388	2.5	6.3	9	Η.	9
Ү869Н9-12	7808-12aa x Y769	7438	3.6	5.7	7	m.	9

TEST 5599. HYBRID PERFORMANCE OF MONOGERM S1 PROGENY LINES, SALINAS, CA., 1999

(cont.)

		Acre Yield	ield		Beets/	Root	
Variety	Description	Sugar	Beets	Sucrose	1001	Rot	RJAP
		sqT	Tons	&	No.	o/e	o(0
S ₁ lines from	S_1 lines from popn-808 (cont.)						
х869н9-13	7808-13aa x Y769	8208	23.68	17.33	168	30.2	87.1
х869н9-16	7808-16aa x Y769	7038	22.44	15.65	135	37.9	86.5
S ₁ lines from popn-818	popn-818						
Y869H15-1B	6818-1Baa x Y769	7552	22.25	17.00	168	25.8	86.2
Y869H15-2B	6818-2Baa x Y769	8407	24.76	16.95	170	33.5	84.5
Y869H15-1	6818-1aa x Y769	7803	23.14	16.85	161	24.5	84.1
X869H15-2	6818-2aa x Y769	7794	23.19	16.80	176	35.2	83.7
X869H15-6	6818-6aa x Y769	7474	22.76	16.40	180	22.6	84.3
Y869H15-21	6818-21aa x Y769	8174	24.22	16.88	136	34.8	85.1
;							
Mean		7712.7	22.92	16.80	169.6	25.9	85.8
LSD (.05)		1407.0	3.99	0.77	26.5	19.0	2.5
C.V. (%)		13.1	12.49	3.29	11.2	52.7	2.1
F value		4.0**	3.33**	4.56**	3.4**	1.7**	1.5*

Roots with obvious rot counted before harvest. Rotted roots Root rot caused considerable variability in plot weighed at harvest but not included in sugar sample. weights. Notes: Root rot due to Sclerotium rolfsii.

TEST 5099. WESTERN SUGAR, BETASEED, & USDA HYBRID EVALUATION UNDER RHIZOMANIA, SALINAS, CA., 1999

Planted: April 29, 1999 Not harvested for yield Scored: October 12-13, 1999 48 entries x 7 reps., RCB 1-row plots, 22 ft. long

	Description	Stand	Count	survi-	Missing Feet	Rot	Koot Kot (Stand)	Knizomania Resistance	Kulzomania Resistance
		Mean	Mean	%	Mean	Mean	o,	Id	8R(0-4)
Western Sugar entries	ntries								
Rizor	rec'd 4-9-99	6	ω	2	•		4.7		6
Crystal 9906		0	о О	9	•		•		9
HM 1646		ö	ω.	6	•		o,		0
Crystal 923R		•	ω.	ω	•		•		4.
Rifle		37.4	29.0	77.5	2.3	7.1	6	3.4	84.6
Beta A943R		0	Η.	0	•	•	•	•	e.
HM 1693		38.7	ω.		1.4	4.7			98.7
Crystal 924R		8	H.	7.	•	•		•	9
Beta 4006R		7.	ÿ.	5	•	•	•	•	7
HM 1645		8	ij	ω.	•	•	7.	•	0
Beta 4038R		41.3	35.6	86.2	1.4	5.0	12.2	2.9	7.96
HM 1647		0	ω.	9	•	•	•	•	6
Beta A940R		7.	w.		•	2.4	•		•
Beta A942R		0	o.	e.	•		⊣.		2
SX Kojak		40.6	28.4	71.1	5.6	7.1	17.0	3.6	74.2
Monohikari		ю Ю	5.	ъ.	•	•	5.	•	0
Western Sugar e	entries (Transgenics)								
Beta 4546LL		•	ω.	Э.	•	5.3	ij.	•	•
Beta A893LL	rec'd 4-9-99	42.1	4.	8	•	6.6		•	٦.
7CG9236LL		ė.	ė.	8	•	•	4.	•	7.
HM 1605RR		æ.	9	8	•	•	o.	•	•
HM RH3RR		÷.	ö	7.	•	•		•	ė.
HM 106RR		ij.	5.	5.	•	•	•	•	ω.
HM 1628RR		9	ω.	5.	•	•	4.	•	0
Crystal 978LL		44.1	32.3	72.9	2.6	8.6	19.8	5.0	28.3
SX0220LL		80	0	0	•	•	9	•	•

TEST 5099. WESTERN SUGAR, BETASEED, & USDA HYBRID EVALUATION UNDER RHIZOMANIA, SALINAS, CA., 1999

(cont.)

Variety	Description	Stand	Harv. Count	Survi- val	Missing Feet	No. Rot	Root Rot (Stand)	Rhizo Resis	Rhizomania Resistance
		Mean	Mean	æ1	Mean	Mean	o-	IO	8R(0-4)
Checks for to	transgenics								
cari	Seedex, 4-16-99, susc. ck.	•		2	•	•	17.5	•	•
Beta 4776R	Betaseed, 1-19-99, resist ck	42.3	36.7	86.9	1.0	5.0	ö.	2.8	98.8
US H11	susc. check	9	•	5.	•	•	16.5	•	•
che che									
SS-432R	Spreckels, 2-8-99	37.4	38.1	100.0	0.7	•	•	•	:
Beta 4430R	Betaseed, 3-10-99	m	38.1	8	1.1	8.0	18.0	5.6	98.3
Beta 4776R	Betaseed, 1-19-99	•	37.7	9	•	•	•	•	7.
Beta 4035R	Betaseed	40.6	34.9	86.3	ნ.0	5.6	•	•	ë.
KWS 6770	Betaseed	•		Η.	•	•	<u>و</u>	•	4
US H11	susc. check	7.	•	•	•	5.7	14.8	•	4
ACC7265	entiles rec'd 3-22-09		_	73.8		10			
1 0 0 0) (•) (•	•
106/2/5		د	۰	'n,	•		'n	٠	7
7KJ5073		о О	,	58.3	•	•	4.	•	83.7
7KY5109		39.9	29.4		2.1	8.7	21.6	3.2	95.1
8CG7343		8	ъ.	84.1	•	•	9	•	93.8
USDA entries R778%H50(Iso)	C790-15CMS x RZM R778		4	о О	3.4				0
R882H27	6831-4HO x	35.3	34.3	97.3		2.1	6.5	3.3	87.3
X869H50	C790-15CMS x Y769 (C69)		9	و	•	7.6	80	•	5
Y869H27	C831-4HO x Y769 (C69)	7.	9	S.	•	•	8		9
X869H5	C833-5aa x Y769 (C69)	ъ.	6	4.	•	4.6	⊢.	•	9
Y875H50 (Iso)		о	m m	.	•		⊢.	•	ω.
8926H50 (Iso)) C790-15CMS x RZM 7926	39.4	25.3	63.8	2.3	11.0	27.6	3.5	83.2
8936H50	x RZM R776-8	6	0	6.	•		9	•	ė.
8939H50	C790-15CMS x RZM Y769H31	o.	o.		•	6.6	4.	3.7	67.1

WESTERN SUGAR, BETASEED, & USDA HYBRID EVALUATION UNDER RHIZOMANIA, SALINAS, CA., 1999 TEST 5099.

(cont.)

		Stand Harv.	Harv.	Survi- 1	Missing	No.	Root Rot	Rhizomania	ania
Variety	Description	Count	Count	Val	בפר	200	(Staild)	DESTS	ance
	21	Mean	Mean	%	Mean	Mean	%	DI %	R(0-4)
Mean	.,	39.8	31.1	78.5	2.1	0.9	15.1	3.7	70.0
LSD (.05)		3.9	7.7	18.8	2.3	4.6	11.3	0.4	10.4
C.V. (%)		9.4	23.7	22.7	102.3	73.1	71.4	9.5	14.1
F value		3.8**	4.3**	4.4*	4.1**	3.3**	3.3**	47.9**	68.4**

the ANOVA for 7 or 8 reps, rep. 5 was deleted. Plots were lifted and layed out by hand. Individual roots that be valid and little affected by rot. Problems occurred in the collection of data for Rep. 5. After reviewing Sugar and root yield data were not collected. Because of the effects of root rot caused by Sclerotium rolfsii, it did not appear that yield data would be These data appear were not severely damaged by rot were scored for rhizomania based on a scale of 0 to 9 where 0 to 4 were meaningful. Roots not infected with S. rolfsii were scored for reaction to rhizomania. considered variations in resistance and 5 to 9, variations in susceptibility. Rhizomania - Entries were evaluated using root scores.

DI = disease index, is the mean score of all roots in an entry. The lower the score, the lesser the visual effects of rhizomania.

resistant vs. susceptible classes did not occur. For example, about 12% of uniformly susceptible US H11 Obviously, the segregation of plants into discrete Likewise, it is likely that some genotypically resistant plants were scored as \$R(0-4) = percentage of plants that appeared to be resistant based on checks and experience with reactions to disease conditioned by the Rz allele. was scored resistant. being susceptible.

previously tested commercial and experimental hybrids. The reaction to rhizomania data appears to have good Overall, the DI and %R values appear to give a good fit to the known relative reactions of the checks and reliability.

		Stand	Harv.	Survi-	Missing	No.	Root Rot	Rhizomania
Variety	Description	Count	Count	val	Feet	Rot	(Stand)	Resistance
		Mean	Mean	જ∘I	Mean	Mean	o% 	DI %R(0-4)

NOTES: (cont.)

Counts were made in an effort to determine if these varieties had differential reaction to S.rolfsii and if host-Sclerotium rolfsii - Southern root rot caused by sclerotium rolfsii occurred throughout this trial. plant resistance could be detected.

Stand count = mean number of plants counted after thinning (prior to canopy closure).

Harvest count = number of roots per plot scored for rhizomania. Does not include roots with S.rolfsii.

No. Rot = number of roots counted at harvest that had root rot, most likely due to S.rolfsii

% Root Rot = percentage of roots with rot at harvest in relationship to initial stand counts

Missing Feet of Plot = measurement of linear feet of row in which rot had destroyed or rotted beets

However, it also rhizomania resistant entries. It may be that resistant, sound roots are more difficult to infect and rot than S.rolfsii was first detected when original stand counts were made. These small plants were quickly destroyed The best indication of sensitivity to S.rolfsii might be the differences between the original stand count and the number of plants that could be scored for rhizomania (% survival). For example, US H11, Monohikari and some of the transgenics appeared to be highly rhizomania impaired roots. Under the conditions of this test there appeared to be a definite differential by rot and disappeared without trace. Infection occurred progressively through out the growing season, (If rhizomania resistant roots are more difficult to infect and rot, seemed that the rhizomania susceptible checks and entries were more susceptible to S.rolfsii than the genetic analysis might suggest that the Rz allele was conditioning partial resistance to S.rolfsii) sensitive to S.rolfsii, whereas Rizor, Beta 4776R et al. appeared to be much less sensitive. usually moving laterally, plant-to-plant within a plot row. varietal reaction to S.rolfsii.

USDA entries - C790-15CMS = rhizomania susceptible, monogerm tester. C78, C82, C69, R76-89-5H31 & Y769H31 segregate for Rz. Y775 & 7926 segregate for resistance from Beta maritima.

TEST 5199. CBGA CODED RHIZOMANIA TEST, SALINAS, CA., 1999

78 entries x 8 reps, RCB 1-row plots, 22 ft. long

Planted: April 29, 1999 Not harvested for yield Scored: October 14-19, 1999

Rhizomania Resistance	<u>%R(0-4)</u>		4	86.5	2	6		S.	81.5	9	8	ω.	2	97.9	2	9.	7.	ö	95.3	금.	w.	თ	8	8	73.0	4.
Rhizo Resis	DI		•	3.3	•	•		•	3.4	•	•	•	•	2.9	•	•	•	•	3.0	•	•	•	•	•	3.5	•
Root Rot (Stand)	≪		•	6.7	•	8.3	8.5	•	6.6	•	12.9			15.7	•		7.5	•	8.6	•	8	H	w.	0	6.3	•
No. Rot	Mean	1.5		5.6	•	3.5		•	3.8	•	•	4.0	•	7.1	•	•	•	•	3.9	•	•	•	•	•	5.6	•
Missing Feet	Mean	•	•	8.0	•	•	1.4	0.4	1.3	9.0	•	•	•	1.1	•	•	•	•	1.4	٠	•	•	•	•	0.4	•
Survi- val	% I	7	ω.	96.1	4.	5	4.	0	9.68	7.	2	6	7.	76.7	m.	σ.	5.	0	100.0	4.	2	4.	9	<u>ი</u>	94.5	80
Harv. Count	Mean	6	0	38.3	9	5.	9	9	33.8	H.	7.	7.	8	34.4	9	5.	5.	4	36.9	4.	7.	5.	9	7.	37.3	9
Stand	Mean		m	39.8	4.	41.3	ω.	5.	37.8	7	•	•	0	44.9	•	o,	7.	5.	37.9	0	1.	7.	8	თ	39.5	0
Source		Spreckels			Spreckels		Spreckels	Spreckels	Spreckels		Spreckels	Spreckels	Spreckels		Spreckels		Spreckels	Betaseed	Betaseed	Betaseed	Spreckels	Spreckels	Betaseed		Spreckels	
Varietv		SS-338R	Beta 4684R	98HX853	98CX20	Beta 4419R	98CX28	Rival	H93203	ഷ		Rodeo	SS-289R	7CG7376	98CX29	Н93392	Pinnacle	4KJ0166	7CG7303	7CG7410	98CX21	SS-781R	5CG7540	7CG7373	99HX913	97CX14
Code No.		SR - 1	- 2	ı m	4	ı U	9	- 7	80	- 9 ¹ Be	-10	-11	-12	-13	-14	-15	-16	-17	-18	-19	-20	-21	-22^{2}	-23	-24	-25

TEST 5199. CBGA CODED RHIZOMANIA TEST, SALINAS, CA., 1999

Code No.	Variety	Source	Stand	Harv. Count	Survi- val	Missing Feet	No. Rot	Root Rot (Stand)	Rhiz Resi	Rhizomania Resistance
			Mean	Mean	∞ 1	Mean	Mean	%	DI	8R(0-4)
SR-26	Rizor	Spreckels	7.	4.	N	•	•	•	•	6
-27	Beta 4488R	Betaseed	41.3	9.		•	6.6	•	•	9.
-28	H945187	Spreckels	9.	ω.	ω	2.3	•	21.4	•	72.4
-29	98CX16	Spreckels	42.6	40.6	95.5	8.0	2.5	•	3.7	72.5
-30	SS-432R	Spreckels	6	5	⊣	6.0	•	8.6	•	61.3
-31	98CX19	Spreckels	9.	4	87.1		•	15.0	•	ω.
-32	98CX857	Spreckels	œ	ω.	0	•	•	•		0
-33	7KJ0146	Betaseed	0	4.	4.	٠	•	•	•	ເດ
-34	Alpine	Spreckels	39.9	37.0	93.5	1.0	2.3	5.5	3.4	81.7
-35	Н92463	Spreckels	7.	,	9	•	•	•	•	7.
-36	98CX30	Spreckels	•	ъ.	80	•		•	•	8
-37	5KJ5057	Betaseed	0	m	4.	•		5.		4.
-38	4KJ0164	Betaseed	2	Η.	5	•	•	•	•	9
-39	99HX915	Spreckels	34.3	26.4	76.8	2.6	8.1	24.1	3.6	74.1
-40	98CX31	Spreckels	0	6	9	•	•	•		5
-41	99HX918	Spreckels	œ.	0	Η.	•	•	16.1	•	8
-42	98CX23	Spreckels	。	ω	9	•	•	•	•	9
-43	98CX32	Spreckels	36.6	34.3	93.8	1.1	3.1	8.6	3.4	0.08
-44	Beta 4210R	Betaseed	9	m.	9	٠	•	•	•	œ
-45	99HX912	Spreckels	œ ·	5.	9	•		4	•	4.
-46	Beta 4300R	Betaseed	2	5.	•	•	12.3	ω.	•	Ξ.
-47	H95555	Spreckels	ω	6	7.	•	•	т М	•	Η.
-48	7KJ0191	Betaseed	5	9	•	٠	•	1.	•	7.
-49	6CG7492	Betaseed	34.6	22.4	66.3	3.3	11.4	32.6	3.3	86.3
-50	SS-778R	Spreckels	ω.	5	•	•	11.5	0	•	7
-51	Rifle	Spreckels	0	4.		•	•	ω.	•	
-52	7CG7408	Betaseed	40.4	27.6	9.79	3.0	0.6	23.0	3.7	68.7
-53	US H11	Standard	0	5.	5	•	•	7.	•	•

TEST 5199. CBGA CODED RHIZOMANIA TEST, SALINAS, CA., 1999

(cont.)

Rhizomania Resistance	DI &R(0-4)	3.2 88.6 2.7 97.7	.5 79. .1 55. .5 77.	4 84 0 93	3.0 96.5 3.3 89.7 2.8 93.3	.0 92. .6 9.	.4 81.	3.9 64.6 3.7 71.8 3.3 84.5	4.8 25.7 2.9 99.7 3.5 78.3 3.3 85.8 4.2 50.9	3.6 74.0 4.0 59.5
Root Rot (Stand)	o∜0	26.3	6.9 0.4.0 9.3		24.4	12.4 19.9	5.7	71.7 9.0 4.4	31.4 10.2 7.4 6.8 17.4	19.0
No. Rot	Mean	11.3 4.8	25.0 8.0 4.		1.9			3.4 1.8	13.0 4.5 3.0 2.6 7.4	3.9
Missing Feet	Mean	4.0	0.1 1.5		0.6	0.5 8.3		n o o. . o. o.	4.1.000.8 0.08 0.1.09	1.5
Survi- val	o∤o	56.8 88.9	90.2 84.4 91.0	731	97.9 63.1 92.4	2 .	900	82.5 91.6 99.3	56.7 86.5 98.3 97.9 78.1	74.5
Harv. Count	Mean	24.1 39.8	35.1 30.4 32.9	6.	41.0 25.4 41.3	N	. 60 60	31.3 34.4 37.4	23.5 35.6 37.6 38.4 32.8	29.4 34.0
Stand	Mean	42.9 44.6	39.1 1.00 1.00 1.00 1.00	9	42.0 41.1 44.5	. 4	2 + 1	37.5 37.5 37.9	41.6 42.0 38.8 39.3 42.3	39.5 39.0
Source		Betaseed Betaseed	Spreckels Spreckels Spreckels	Spreckels Betaseed	Betaseed Betaseed Betaseed	Betaseed Betaseed	Spreckels Spreckels	Spreckels Spreckels Spreckels	Betaseed Betaseed Spreckels Spreckels	Spreckels Spreckels
Varietv		7CG7321 5KJ0142	99HX914 98CX861 Imperial	imperiar SS-NB7R 6KJ0163		Beta 4035R 7CG7621	Summit 97CX01	99HX917 98CX858 Phoenix	8CG7064 7CG7322 99HX926 99HX928	99HX925 99HX924 eck
Code		SR-54 -55	- 56 - 57 - 57	65-1 69-1	-61 -62 -63	-64 -65	99-	-68 -69 -70	-71 -72 -73 -74	-76 9 -77 9 USDA check

TEST 5199. CBGA CODED RHIZOMANIA TEST, SALINAS, CA., 1999

Code No.	Variety	Source	Stand	Harv. S	Survi- val	Missing Feet	No. Rot	Root Rot (Stand)	Rhizomania Resistance	ania cance
			Mean	Mean	ø₽ 		Mean	o⁄o l	Id	SR (0-4)
Mean			39.5	33.1	84.3	1.6	5.6	14.2	3.5	6.94
LSD (.05)			4.1	6.9	16.2	1.8	3.9	8.6	0.3	9.4
C.V. (%)			10.5	21.1	19.5	19.5 117.8	71.9	70.8	7.8	7.8 12.4
F value			3.3**	5.0**	4.8**	4.2**	4.4*	4.7**	31.9**	33.0**

Entry 9 had 0% emergence. Beta 4776R transplants used as filler. Entry 22 had low frequency of bolters (annuals)

meaningful. Roots not infected with S. rolfsii were scored for reaction to rhizomania. These data appear to be valid and little affected by rot. Plots were lifted and layed out by hand. Individual roots that were not NOTES: Rhizomania - Entries were evaluated using root scores. Sugar and root yield data were not collected. Because of the effects of root rot caused by Sclerotium rolfsii, it did not appear that yield data would be severely damaged by rot were scored for rhizomania based on a scale of 0 to 9 where 0 to 4 were considered variations in resistance and 5 to 9, variations in susceptibility.

DI = disease index, is the mean score of all roots in an entry. The lower the score, the lesser the visual effects of rhizomania.

resistant vs. susceptible classes did not occur. For example, about 12% of uniformly susceptible US H11 reactions to disease conditioned by the Rz allele. Obviously, the segregation of plants into discrete was scored resistant. Likewise, it is likely that some genotypically resistant plants were scored as %R(0-4) = percentage of plants that appeared to be resistant based on checks and experience with being susceptible.

The reaction to rhizomania data appears to have good Overall, the DI and &R values appear to give a good fit to the known relative reactions of the checks and previously tested commercial and experimental hybrids. reliability

CBGA CODED RHIZOMANIA TEST, SALINAS, CA., 1999 TEST 5199.

Rhizomania	lesistance	&R(0-4)
R.	Re	DI
Root Rot	(Stand)	o⊱
No.	Rot	Mean
Missing	Feet	Mean
Survi-	val	op]
Harv.	Count	Mean
Stand	Count	Mean
	Source	
	Variety	
Code	No.	

NOTES: (cont.)

were made in an effort to determine if these varieties had differential reaction to S.rolfsii and if host-Sclerotium rolfsii - Southern root rot caused by sclerotium rolfsii occurred throughout this trial. plant resistance could be detected.

Stand count = mean number of plants counted after thinning (prior to canopy closure).

Harvest count = number of roots per plot scored for rhizomania.

Does not include roots with S.rolfsii.

No. Rot = number of roots counted at harvest that had root rot, most likely due to S.rolfsii.

% Root Rot = percentage of roots with rot at harvest in relationship to initial stand counts.

Missing Feet of Plot = measurement of linear feet of row in which rot had destroyed or rotted beets.

However, it rhizomania resistant entries. It may be that resistant, sound roots are more difficult to infect and rot than These small plants were quickly destroyed usually moving laterally, plant-to-plant within a plot row. The best indication of sensitivity to S.rolfsii might be the differences between the original stand count and the number of plants that could be scored for rhizomania (% survival). For example, US H11 and some of the more susceptible entries appeared to be highly also seemed that the rhizomania susceptible checks and entries were more susceptible to S.rolfsii than the rhizomania impaired roots. Under the conditions of this test there appeared to be a definite differential by rot and disappeared without trace. Infection occurred progressively through out the growing season, (If rhizomania resistant roots are more difficult to infect and rot, genetic analysis might suggest that the $R\mathbf{z}$ allele was conditioning partial resistance to S.rolfsii) . sensitive to S.rolfsii, whereas Beta 4430R, Beta 4776R et al. appeared to be much less sensitive. S.rolfsii was first detected when original stand counts were made. varietal reaction to S.rolfsii.

32 entries x 8 replications, RCB(E) 1-row plots, 27 ft. long (16 blocks, 16 rows)

Planted: September 23, 1998 Harvested: June 15, 1999

		Acre Yield	ield		Beets/		Clean	
Variety	Description	Sugar	Beets	Sucrose	1001	Bolters	Beets	NO3-N
		I.bs	Tons	o(o	No.	%	o(○	Mean
Checks R4776R	Beta 4776R 7653 (3-27-98)	10607	77	4.6	155	0.0		188
Rifle	. L11624	11016			149	•	4	190
E-000000000000000000000000000000000000	1206/20MS							
1	4807HO (C306/2CMS) x Y7	10984	42.64	12.87	137	0.3	94.8	211
R882H37	4807HO (C306/2CMS) x R781,R776	9330	39.25	11.82	139	0.0	95.1	223
Testcrosses to	C790-15CMS							
R576-89-18H50	C790-15CMS x R476-89-18	12106	40.65	14.90	150	2.1		143
R876-89-5NBH501 C790-15CMS	x RZM-% R576-89-	10081^{1}	•	15.29	136	1.3	93.0	111
R876-89-5H50	C790-15CMS x RZM-% R576-89-5	12305	41.62	14.77	154	9.0	94.0	133
R882H50	C790-15CMS x R781,R776	10460	36.21	14.43	140	0.3	94.4	148
Y869H50	C790-15CMs x Y769 (C69)	11658	σ.	14.58	142	1.6	94.2	174
Y868H50	C790-15CMS x RZM Y768	11083	38.83	٦.	145	0.0	94.3	171
R878H50	C790-15CMS x R778, R778% (C78)	160	۲.	4.	133	1.7	•	147
R854H50	C790-15CMS x RZM R754	10902	38.51	4.1	4	0.7	93.3	148
Y873BH50	C790-15CMS x RZM Y773	10616	37.62	14.07	147	1.6	94.1	187
Y875H50	C790-15CMS x RZM Y775,	10457	8	4.1	141	1.0	93.4	172
X866H50	C790-15CMS x RZM Y766	11505	ъ В	٠.	147	•	94.4	172
X867H50	C790-15CMS x RZM Y767 (C67)	10386	37.64	13.84	148	9.0	94.4	183
X871H50	C790-15CMS x RZM Y771	12937	44.31	14.62	151	1.5	5	128
X872BH50	C790-15CMS x RZM Y772 (C72)	12216	3.0	4.1	147	0.0	ж Э	161
X872H50	C790-15CMS x RZM-% Y672	12913	45.30		160	2.6	93.5	188
8931H50	C790-15CMS x RZM 7931	11890	. 2	4.3	145	0.3	m.	160

EVALUATION OF TESTCROSS HYBRIDS, IMPERIAL VALLEY, CA., 1998-99 TEST B199.

(cont.)

		Acre Yield	eld		Beets/		Clean	
Variety	Description	Sugar	Beets	Sucrose	1001	Bolters	Beets	NO3-N
		Lbs	Tons	o(0	No.	op	<i></i> %∣	Mean
Testcrosses t	to C790-15CMS (cont.)							
8924H50	C790-15CMS x RZM 7924	11569	41.10	14.05	144	0.7	93.7	159
8932H50	C790-15CMS x 7932CT,7201	10793	37.87	14.37	140	2.8	92.6	186
2831H50	C790-15CMS x RZM Z730, Z731	11446	41.38	13.81	144	0.3	93.2	168
8926H50	C790-15CMS x RZM 7926	11309	37.72	15.04	155	1.3	92.5	116
	01-0100 9-00-Mad : 0200-1-0000	12832	24 23	14.46	150	8.0	93.5	183
8913-1000	C/ ACTOMA A NAM-BN-8 COLO /0	100			977			000
8935H50	C790-15CMS x R776-89-5H13	12560	47.94	T4.60	14p	٥. ٤	γ) Υ)	139
8936H50	C790-15CMS x RZM R776-89-5H31	11222	37.60	14.96	139	0.7	92.8	131
8937H50	C790-15CMS x RZM R776-89-5H11	12041	40.10	15.03	150	0.0	93.9	117
8938H50	C790-15CMS x RZM Z731H11	12218	40.87	15.01	157	1.0	94.3	129
8939450	C790-15CMS x RZM Y769H31	11569	40.16	14.46	145	1.6	93.0	124
8918-12H50	C790-15CMS x RZM-ER-% 6918-12	12325	43.20	14.29	126	0.3	95.2	164
8918-21H50	C790-15CMS x RZM 7918-21	9785	31.97	15.28	127	0.3	93.4	109
		1	0	•			0	0
Mean		11397.8	39.62	T4.40	144.y	ا س	y3.8	7.001
LSD (.05)		1298.8	3,83	0.78	11.1	1.7	1.5	56.0
C.V. (%)		11.6	9.80	5.47	7.8	195.8	1.6	36.0
F value		3.9**	5.09**	5.90**	4.0**	t 1.6NS	2.0**	2.1*

4aa x CZ25), and Y769H31 = F_1 (popn-931aa x C69) are F_1 population and line hybrids that are being evaluated and improvement in the breeding program at Salinas. Lines R754, Y766, Y767, Y771, Y772, Y773, Y775, and 7926 have 89-5), R776-89-5H31 = F_1 (popn-931aa x C76-89-5), R776-89-5H11 = F_1 (C911-4aa x C76-89-5), Z731H11 = F_1 (C911resistance to rhizomania and at Salinas for nonbolting, virus yellows resistance, performance per se, and for resistance to rhizomania and germplasm from Beta vulgaris ssp. maritima. R776-89-5H13 = F_1 (C913-70aa x C76yellows, bolting, etc. S1 progenies from these F1 hybrids are being evaluated at Brawley for nonbolting and Individual plants within these F1 hybrids should be Aa, Sf, and segregate for resistance to rhizomania, virus NOTE: The pollinators of these experimental hybrids are breeding lines and populations under population developed as potential sources for S_1 progeny selection for combined disease resistance and performance. resistance to rhizomania, powdery mildew, and Erwinia.

¹R876-89-5NBH50 had appearance of C76-89-5 pollinator and not its H50 hybrid. Pollinator seed may have been planted

EVALUATION OF EXPERIMENTAL HYBRIDS (POPN & s_1 PROGENY TESTCROSSES), IMPERIAL VALLEY, CA., 1998-99 TEST B399.

Planted: September 23, 1998 Harvested: May 17, 1999¹ 32 entries x 8 reps., RCB(e), 2 blocks per rep 1-row plots, 27 ft. long, 16 blocks, 16 rows

		Acre	Acre Yield		Beets/		Root	Clean	
$Variety^2$	Description ²	Sugar	Beets	Sucrose	1001	Bolters	Rot	Beets ³	NO3-N
		Lbs	Tons	o⊱	No.	æ	æ	%	Mean
Checks		7		r L	((ć
Rizor	- 6 - 6 - 3 -	12131	טי	ກ .	י פ	•	•	د	ָ מ נ
SS-778R	9-98,	11887	1.1	4.4	ဖ	٠	٠		102
Rifle	-	11859	38.97	15.18	165	2.1	0.0	92.1	107
B4776R	Beta 4776R.7653 (3-27-98)	11034	5.7	5.4	9	•	•	7	96
R878H50	C790-15CMS x R778, R778%	_	0.1	4.3	Ω	•		8	H
R882H50	C790-15CMS x R781, R776,	10876	38.02	14.31	155	0.0	0.7	92.8	116
Population hy	hrids								
8931H50 C790	C790-15CMS x RZM 7931	11693	39.85		149	0.3	0.4	92.1	93
8924H50	C790-15CMS x RZM 7924	035	5.8	4.5	Ω	2.6	•	;	122
מינון מינוטטטטטטטטטטטטטטטטטטטטטטטטטטטטטטטטטטטט	ت. بازی								
8013-70450 6700	C700-15CMS * B7M-EB-* 6013-70	9788	o v	٨	165	ζ.		_	100
8911-4-10H50)-15CMS × BZM-ER-%	008		י ה	163	•	•	, –	1 (
8918-12H50	x RZM-ER-% 6918-12	10975	•	14.36	143	0.1		0.06	94
8918-21H50	× RZM 7918-	001	2.6	5.3	ϵ	•	0.0	9	92
8925-19H50	C790-15CMS x 6925-19	233	2	4.0	160	•	0.0		105
8929-41H50	$C790-15CMS \times 6929-41$	12219	0		162	0.0	0.0	88.0	77
8929-72H50	$C790-15CMS \times 6929-72$	121	8	5.2	157	•	•	Ξ.	64
8929-102H50	C790-15CMS x 6929-102	225	1.7	4.6	Ŋ	•	0.0	m.	103
8929-112H50	C790-15CMS x 6929-112	150	7.9	5.1	158	•	•		81
8929-114H50	$C790-15CMS \times 6929-114$	30	8.1	5.4	\mathbf{S}	•	0.3	0	57
8929-115H50	$C790-15CMS \times 6929-115$	10919	35.39	15.47	147	1.5	0.0	89.5	97
8929-133H50	C790-15CMS x 6929-133	7.1	1.4	5.6	$^{\circ}$	•	0.0	5.	61

EVALUATION OF EXPERIMENTAL HYBRIDS (POPN ϵ s_1 PROGENY TESTCROSSES), IMPERIAL VALLEY, CA., 1998-99 TEST B399.

(cont.)

		Acre Yield	ield		Beets/		Root	Clean	
$Variety^2$	Description ²	Sugar	Beets	Sucrose	1001	Bolters	Rot	Beets ³	NO3-N
		Irbs	Tons	₩	No.	%	φI	ا% ا	Mean
S ₁ Progeny Hy	S ₁ Progeny Hybrids (cont.)								
8930-19H50	C790-15CMS x 6930-19	12582	41.95	14.96	158	0.0	0.0	90.2	84
8930-39H50	C790-15CMS x 6930-39	12030	40.57	14.85	161	0.0	0.0	7.06	80
8930-102H50	$C790-15CMS \times 6930-102$	10693	35.09	15.24	164	0.0	0.0	98.6	106
Z825-6H50	C790-15CMS x Z625-6	12388	41.41	14.99	167	1.9	0.0	8.68	82
Z825-9H50	C790-15CMS x Z625-9	10829	34.79	15.60	152	0.3	0.3	90.2	83
Z830-11H50	$C790-15CMS \times Z630-11$	13284	45.80	14.48	160	5.6	0.0	91.5	06
8927-29H50	$C790-15CMS \times 6927-29$	11030	36.21	15.28	155	2.3	0.0		68
8927-30H50	C790-15CMS x 6927-30	11372	36.99	15.35	170	0.8	0.0	86.3	57
8927-33H50	C790-15CMS x 6927-33	9280	30.50	15.26	160	5.6	0.0	88.5	83
8927-37H50	$C790-15CMS \times 6927-37$	11137	38.55	14.46	157	16.8	0.0	91.5	115
8929-153H50	$C790-15CMS \times 6929-153$	11287	38.38	14.79	159	0.0	0.0	92.4	78
8929-154H50	$C790-15CMS \times 6929-154$	10593	37.55	14.11	158	0.0	0.0	89.1	127
Mean		11178.1	37.61	14.90	156.9	1.9	0.1	90.1	89.2
LSD (.05)		1297.9	3.84	0.87	12.2	3.5	0.5	2.2	40.0
C.V. (%)		11.8	10.37	5.92	7.9	182.8	878.7	2.5	45.5
F value		e.6**	10.23**	2.95**	3.5**	6.9 **	0.9NS	**8.8	2.3**

¹Harvested 11 days after last irrigation under moderately wet conditions and high fertility.

S₁'s were selected at Salinas on basis of per se disease resistance and performance and testcrossed R776 = C82. 7931 & 7924 = MM,S^f,A:aa popns. S₁ progeny were individually selfed plants from 2 R778, R778% = C78. MM, S^f, A: aa popns. to C790-15CMS.

³See Test B499.

16 entries x 8 reps., RCB(E), 2 blocks/rep 1-row plots, 24 + 3 ft. long, 16 blocks, 8 rows

Planted: September 23, 1998 Harvested: May 14, 1999

			•					į.	
Variety	Description ²	Acre Y Sugar	Yield Beets	Sucrose	Beets/ 100'	Bolters	Root Rot	Root Clean Rot Beets	NO3-N
		I.bs	Tons	%	No.	o(∙	%	%	Mean
Checks	0+12 1776 7653 (3-27-08)	10572	25 77	α	169	o o	C	ور ر	162
X869H50	C790-15CMS x Y769	693	6.0	m	161			92.1	202
X869H37	4807HO (C306/2CMS) x Y769	9230	36.70	2	156	0.0	0.0	92.8	217
Population hybrids	hybrids								
Y869H35	7835mmaa x Y769	9105	4	•	147	•	•	4	210
х869нзв	7838mmaa x Y769	9702	35.72	13.57	148	0.3	1.9	94.7	151
	hybrids								
X869H7	6911-4-7HO x Y769	9466	6.4	•	147	0.0	9.0	92.3	156
Y869H45	7867-1HO x Y769	σ	35.03	12.81	147	6.0	6.0	91.0	198
X869H46	7869-6HO x Y769	9181	5.4	12.97	157	9.0	9.0	91.7	158
Y869H27	6831-4HO x Y769	10156	38.61	13.16	151	9.0	2.6	94.0	167
Y869H4	5831-3aa x Y769	7456	28.74	13.07	129	0.0	8.4	95.0	161
X869H5	5833-5aa x Y769	9468	Т.	•	154	•	•	92.9	142
X869H12	5833-12aa x Y769	7917	30.44		147	0.3	7.2	5.	180
X869H29	5829-3aa x Y769	7769	28.17	13.78	155	0.0	2.5	93.4	139
X869H15-2B	6818-2Baa x Y769	8028	0.7	13.14	150	0.0	2.4	ij	165
X869H15-6	6818-6aa x Y769	7696	28.55	13.48	153	•	8.6	95.3	157
X869H15-21	6818-21aa x Y769	7050	5.6	13.80	123	0.0	13.4	•	159
Mean		80.	•	13.41		0.2	3.4	93.3	170.2
LSD (.05)		0.006	3.27	0.71	11.2	•	4.6	1.8	51.6
C.V. (%)		0	6	ω.		ن	136.2	•	30.6
F value		11.2*	* 10.78**	4.58**	7.6**	1.5NS	6.0**	5.4**	1.7NS

	NO3-N	Mean
Root¹ Clean	Beets	æ1
$Root^1$	Rot	90
	Bolters	%
Beets/	1001	No.
	Sucrose	%
Acre Yield	Beets	Tons
Acre	Sugar	rps
	Description ²	
	Variety	

beet may show only superficial rot or may eventually completely rot and die. Of interest to breeders is the distinct susceptible. Multigerm lines appear to be variable in their reaction. Experimental hybrids may be completely free of se were evaluated at Brawley. Resistance to LIYV was the primary objective at the time but it was observed that wide differences occurred for reaction to the black forming crown/root rot. C790-15 was selected partially based upon its high resistance to this root rot. It is not known if this root rot is important in commercial fields in the Imperial problem. Counts in this test were made at harvest and under a full canopy. Actual incidence of this rot is probably crown/root rot or may have a significant number of infected plants. In developing C790-15, for example, S1 lines per or at more advanced stages. If in the seeding stage, the major problems is loss of stand. If in larger beets, the crown/root rot is Rhizoctonia, although Phoma has also been suspected. Plants may be infected in the seeding stage discoloration (jet black skin) to deeper, dry lesions, crown rot, root splitting, destruction of shoot (crown) and subsequent loss of root. During harvest, affected plants often break and the lower tap root is left in the ground. Valley, where I have rarely seen it, or just in the field plot areas on the IDRS. Because of the availability of resistance to this disease, it may be that in the trialing process by researchers, seed companies, processors and genetic diversity and differential reactions among sugarbeet breeding lines and hybrids. Lines C562, C301, C306, growers, that the most susceptible materials do not yield well enough to be retained in the tests. At the very least, based upon observations within tests on the IDRS, Brawley, it will be wise to be aware of this potential Infected plants often occur in multiples of 2 or more down a plot row. The best guess as to the cause of this crown/root rot. This malady is characterized by a crown and upper root infection that leads from superficial 1 For many harvests on the IDRS, Brawley, it has been observed that differential reactions occur to an unknown C790-15, for example, appear to be mostly resistant. Many other monogerms appear to be somewhat to fully much higher.

6911-4-7HO = C911-4-7CMS. 7867-1HO = C867-1CMS. 6831-4HO = C831-4CMS. 5831-3 = C831-3. 5829-3 = C829-35833-12 = C833-12. 2 Y769 = C69.

³Test was harvested very wet (8 days post irrigation) and under high nitrogen status.

Planted: September 23, 1998 Harvested: June 11, 1999

32 entries x 8 reps, RCB(E) 1-row plots, 27 ft. long

, , , , , , , , , , , , , , , , , , ,	n::))								
			Acre Yield	Yield		Beets/		Clean	
Code	Variety	Source	Sugar	Beets	Sucrose	1001	Bolters	Beets	NO3-N
			Ths	Tons	₩	No.	₩	oo I	Mean
98M -23	Beta 4430R	Betaseed	73	5.4	6.2	9	•	4.	06
-25	8CG7064	Betaseed	330		5.3	9	•	5	88
-22	7CG7322	Betaseed	344	4.4	5.1	4	•	4.	ω
-21	Summit	Spreckels	13011	44.61	14.56	148	0.3	93.9	117
1 1	7CG7321	Betaseed	287	1.9	5.3	9	•	ij.	⊣
-18	SS-778R	Spreckels	273	7.	5.0	Ω	•	4	
- 4	Beta 4035R	Betaseed	271	1.5	5.2	2	•	4.	$^{\circ}$
- 2	Rizor	Spreckels	243	9.2	5.8	5	•	æ.	2
-26	7KJ0191	Betaseed	12316	37.28	16.57	156	0.3	93.7	129
ი I	Phoenix	Spreckels	207	1.7	4.4	9	•	5.	S
-13	Rifle	Spreckels	166	6.4	5.9	2		5.	ന
-10	Alpine	Spreckels	69	9.7	4.6	S	•	4	8
-27	Beta 4776R	Betaseed	133	5.8	5.8	Ŋ		4.	2
9 -	7CG7400	Betaseed	П	37.59	15.25	151	0.3	94.5	116
-19	98HX853	Spreckels	31	7.6	5.0	4	•	4.	₩.
-15	Beta 4684R	Betaseed	117	5.4	5.7	9	1.4	5	0
-11	Beta 4210R	Betaseed	169	3.3	3.5	2	•	9	2
-12	SS-NB7R	Spreckels	128	8.0	4.8	9	•	5.	$^{\circ}$
-14	Pinnacle	Spreckels	11353	40.64	13.94	150	4.1	94.8	169
-16	SS-781R	Spreckels	109	8.6	4.3	2	•	5	0
ا 5	5CG7540	Betaseed	102	7.2	4.8	9		2	4
- 7	97CX01	Spreckels	11153	38.33	14.52	160	o.0	94.2	118
& I	Rival	Spreckels	056	4.3	5.3	Ŋ		5.	$^{\circ}$
-17	Imperial	Spreckels	035	7.3	3.8	4		5	ω
-28	US H11	Standard	02	7.7	4.4	ന		2	\leftarrow

TEST B299. AREA 5 CODED NON-RHIZOMANIA YIELD TEST, IMPERIAL VALLEY, CA., 1998-99

(cont.)

			Acre Yield	ield		Beets/		Clean	
Code	Variety	Source	Sugar	Beets	Sucrose	1001	Bolters	Beets	N03-N
			I.bs	Tons	%	No.	o⊱	o⇔	Mean
USDA entries	-972 * (8MD6/908D) -975-	7-88-9-8	12301	42 79	7 7 7 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8	ر م	o C	u 5	7
_	7.000-15.000 x C76-89-5)	11786	37.66	15.67	1 1 2 6	о и О <		0 7 0
	4807HO (C306/2CMS) x C82	x C82	11936	42.90	13.88	147	#	96.0	138
	C790-15CMS x C78		11325	37.46	15.11	154	8 9.5	94.5	106
		C C	•	6	•	,			
R778H37	4807HO (C306/2CMS) x C78	x C78	11344	40.06	14.14	148	1.3	95.5	117
R882H27 (C831-4HO x C82		11139	39.48	14.13	141	2.2	95.7	120
R882H50	C790-15CMS x C82		10727	36.18	14.86	145	3.2	95.3	120
Mean			11731.8	39.27	14.94	153.5	3.7	94.5	127.9
LSD (.05)			1206.4	3.82	0.54	11.1	3.5	1.1	39.4
C.V. (%)			10.4	68.6	3.66	7.4	94.9	1.1	31.3
F value			7.17*	7.17** 6.98**	14.95**	2.6NS	28.3**	6.8**	5.7

(cont.)

Impur.	Value	8060	8599	10232	10341	10053	9931	10169	9290	9143	10339	10395	9801	8971	11434	10063	10002	12356	10841	11021	9951	10711	11679	9712	10898	10
NH2-N	udd	279	312	351	352	364	333	415	354	363	413	444	347	350	452	354	391	441	434	369	363	458	437	355	363	446
Potassium	шdd	1743	1648	2156	2194	2082	2121	2000	1879	1805	2017	1907	2090	1751	2305	2176	1912	2453	2126	2259	2026	ത	ന	1991	\sim	19
Sodium	wdd	301	432	431	433	397	\vdash	S	Ŋ	338	O	403	9	361	394	9	430	582	402	533	410		439	388	522	385
Known SugarLoss	<u>lbs/a</u>	1097	1111	1353	1376	1259	25	1272	1096	1044	1272	1132	1162	973	1276	1133	1055	1595	1234	1343	1151	19	1341	979	$^{\circ}$	932
Recover. Sugar	d⊘	92.5	1.	•	89.3	•	90.1	0.06	91.2	•	89.2	90.2	6	91.4		89.9	90.4	86.1	89.0	ω.	89.5	89.1	7.	90.5	œ.	88.5
Recover. Sugar	1bs/t	300	8	272	260	276	272	275	289	304	258	289	264	289	271	271	285	233	264	246	257	265	255	278		255
Recover. Sugar	<u>lbs/a</u>	13633	12192	12096	11635	11612	11478	11440	11335	11273	10807	10535	10533	10362	10213	10178	10120	10102	10055	10010	9946	9826	9812	9583	9125	7089
Variety		Beta 4430R	8CG7064	7CG7322	Summit	7CG7321	SS-778R	Beta 4035R	Rizor	7KJ0191	Phoenix	Rifle	Alpine	Beta 4776R	7CG7400	98HX853	Beta 4684R	Beta 4210R	SS-NB7R	Pinnacle	SS-781R	5CG7540	97CX01	Rival	Imperial	US H11
Code		98M -23	-25	-22	-21	П	-18	- 4	- 2	-26	ი 	-13	-10	-27	9 1	-19	-15	-11	-12	-14	-16	I J	- 7	8	-17	-28

AREA 5 CODED NON-RHIZOMANIA YIELD TEST, IMPERIAL VALLEY, CA., 1998-99 TEST B299.

(cont.)

	امد		0	0	. ^	ı m	ď	ח מ	റരാ		13.1	1558 O	. L	2.7**
Impur	Value		10410	9939	10632	10113	11733	11 400	10008		10293 1	ا ا ا	, , , ,	* * • M
NH ₂ -N	udd		367	400	358	379	421	110	435		385.0	82.5	α το	2.3
Potassium	udd		2146	1836	2302	2082	2507	2454	1882		2082.7	320.2	15.6	3.6**
Sodium	wdd		446	443	422	373	419	391	335		408.3	119.9	29.8	1.8*
Known SugarLoss	lbs/a	,	1332	1128	1369	1138	1412	1364	1089		1209.4	224.7	18.9	3.3**
Recover. Sugar	o⊳	(89.1	90.4	88.5	6.68	87.5	87.8	6.68	,	9.68	1.7	2.0	4.8**
Recover. Sugar	lbs/t	((422	284	246	272	248	248	267		267.9	12.8	4.8	13.9**
Recover. Sugar	<u>lbs/a</u>	0	6960T	10658	10567	10187	9931	9775	9638	1	10522.4	1117.9	10.8	4*6.7
Variety		210	,	20										
Code		USDA entries	. HC - 60 - 61 - 71	R876-89-5H50	R882H37	R878H50	R778H37	R882H27	R882H50		Mean	LSD (.05)	C.V. (%)	F value

Entries 23 (tall canopy) and 25 (short canopy) were yellowish Powdery mildew was controlled with sulfur. and appeared to be infected with LCV (lettuce chlorosis virus). NOTES: Test was under fairly high nitrogen conditions. appeared to be no significant disease or pest problems.

Planted: September 23, 1998 Harvested: May 13, 1999

48 entries x 8 reps., RCB(E), 3 blocks per rep 1-row plots, 18 ft. long, 24 blocks, 16 rows

		Acre	Yield		Beets/		Root	Clean	
Variety	Description	Sugar	Beets	Sucrose	1001	Bolters	Rot	Beets	NO3-N
		sqT	Tons	%	No.	%	₩	o40	Mean
Checks									
B4776R	Beta 4776R.7653 (3-27-98)	8439	0.6	4.6	വ	0.0	•	0	σ
Rifle	Spreckels, 9-98, L1162401	8074	7.6	4.5	$^{\circ}$	•		6	7
SS-778R		7178	28.26	12.76	145	0.0	0.0	88.9	247
4035R	Betaseed, 7-10-97	7605	9.6	2.8	4	0.0	•	0	7
Testcrosses to	C306/2CMS								
R776-89-5H37	4807HO (C306/2CMS) x R576-89-5	85	7.5	ი.	139	0.0	0.0		ω
R778H37	4807HO (C306/2CMS) x R678	6332	26.06	12.18	136	•	0.0	88.4	205
8926н37	4807HO (C306/2CMS) x RZM 7926	36	1.7	11.67	$\boldsymbol{\omega}$	0.0	0.0	89	ω
+ 0000CTC+00E	2790-1508s								
R882H50		7695	30.16	12.78	139	0.0	0.0	90.2	259
R878450(Sp)	C790-15CMS * R778 R778%	7542	σ	ď	145	1 4			71 م
78, 5515, 551 Y869H50	6977 ×	7442	, 6	9 6	7	•	•		1 0
R876-89-5NBH50	x RZM-%	0909		14.57	133		1.6	80.6	124
Y873BH50	C790-15CMS * RZM Y773	6568	6.3	2.5	$^{\circ}$	•	•	9	7
X867H50	C790-15CMS x RZM Y767 (C67)	7328	8.4	3.1	ന	1.0	0.0	ω.	0
X872H50	C790-15CMS x RZM-% Y672(C72)	8026	1.6		4	•	•	•	Н
Y875H50 (Iso)	×	6283	24.95		133	•	0.0	87.3	266
Y875H50 (Sp)	C790-15CMS x RZM Y775,	9889	7.6		m	0.4	0.0	•	Ŋ
CR813H50	C790-15CMS x RZM CR713	48	3.7	2.6	4	•	•	0	~
8931H50 (Sp)	C790-15CMS x RZM 7931	œ	0.5	2.8	4	0.0	٠	ω	Н
8924H50	x RZM	7526	27.88	13.51	138	0.0	0.0	88.9	157
Z831H50	C790-15CMS x RZM Z730, Z731	9	0.1	2.8	4	0.0		о	m

TEST B699. EVALUATION OF EXPERIMENTAL HYRIDS UNDER RHIZOMANIA, IMPERIAL VALLEY, CA., 1998-99

		9	Yield		Beets/		Root	Clean	
Variety	Description	Sugar	Beets	Sucrose	1001	Bolters	Rot	Beets	NO3-N
		Tps	Tons	ov∘	No.	ok∘	o(0	ا%	Mean
Testcrosses to	C790-15CMS (cont.)								
8935H50 (Iso)	C790-15CMS x RZM R776-89-5H13	7467	7.7	3.5	144	•		ω.	2
8932H50	C790-15CMS x 7932CT,7201	5982	2.5	3.4	140	•	•	5.	7
8926H50 (Iso)	C790-15CMS x RZM 7926	7627	29.24	13.03	154	0.4	1.9	86.5	196
8926H50 (Sp)	C790-15CMS x RZM 7926	7693	9.6	2.9	141	•	•	7.	g
Testcrosses to	o C831-4HO								
R882H27	6831-4HO x R781,R776	7466	1.1		131	•		2	261
X869H27	6831-4HO x Y769	7595	30.54	12.48	140	0.5	0.0	89.9	252
X875H27	6831-4HO x RZM Y775	7023	9.4	11.93	142	•	•	•	228
Testamosses to	698-ngog 3 58-ngog, 888-ngog c								
R882H38	x R781, R776	7371	27.63	13.38	129	0.0	0.0	0.68	147
хв69нзв	7838mmaa x Y769	7758	0.1	σ.	134	•	•	∺.	223
X869H35	7835mmaa x Y769	7085	7.3	3.0	152	0.0	•	ä	247
хв69н69	7869aa x Y769	6819	26.16	13.17	149	0.4	0.0	90.2	218
R878H69	7869aa x R778,R778%	7909	9.7	3.3	149	0.7	•	6.	195
8931H38	7838mmaa x RZM 7931	7533	9.1	σ.	147	•		œ	0
8935H38	7838mmaa x R $776-89-5$ H	7032	7.22	2.9	145	•	•	8	8
8932H38	7838mmaa x 7932CT,720	6490	26.281	12.49	145	0.0	0.0	98.6	217
R878H55	7835H50 x R778,R778%	7721	9.8	2.9	143	•	•	÷.	m
R878H58	7838H50 x R778,R778%	7904	9.6	13.31	4	•		6	177
8931H55	7835H50 x RZM 7931	7528	0.0	2.6	149	•	•	6	267
8931H58	7838H50 x RZM 7931	7599	29.99	12.82	135	0.5	0.0	88.2	204
8926H55	7835H50 x RZM 7926	7650	0.1	. 7	m	•	•	0.	265
8926H58	7838H50 x RZM 7926	7981	31.96	12.48	141	0.5	0.0	88.4	231

(cont.)

		Acre Yield	ield		Beets/		Root	Clean	
	מסייד לידים מסר	Sugar	ts	Sucrose	1001	Bolters	Rot	Beets	NO3-N
Variety		Ibs	Tons	olo	No.	o(0	%I	બ∘	Mean
Topcrossed with C69	th C69	5471	22.13	12.44	120	0.0	8.3	91.4	203
Y869H4	5651-344 X 1/67	5972	23.05	12.98	129	0.0	2.9	91.0	198
хвеянэ Y 869H12	5833-12aa x Y76	6754	25.639	13.21	133	0.5	0.0	92.0	199
9		5973	77.77	13.19	148	0.0	0.0	6.68	148
Y869H29	5829-388 X 1/69	2.00	27 02	12.60	140	1.9	0.0	91.1	265
X869H45	89/IX OHT-/98/	0400	25.75 25.46	12.69	148	0.8	0.5	87.7	212
Y869H46	7869-6HO x 1/69	200	0 1 1			0	c	0	242
X869H7	$6911-4-7H0 \times Y7$	6495	26.8169	12.29	170	o))))	1
		7215.6	28.13	12.90	140.7	0.4	0.3	89.1	221.2
Mean		9.050	3.31	0.75	15.5	1.2	2.1	3.2	71.7
LSD (.05)		13.1	11.94	5.93	11.2	317.8	655.4	3.7	32.9
C.V. (%) F value		4.3**		4.61**	1.7NS	1.4NS	3.1**	2.8**	2.4NS

infected with lettuce chlorosis virus (LCV) and BWYV although symptoms were mostly masked by the high fertility level. - Sugarbeet cyst nematodes were observed at harvest. Powdery mildew was controlled with sulfur. Plants were probably performance relative to tests in Field K without rhizomania suggested rhizomania was an important factor in yield. Harvest under moist, high fertility conditions. Roots at harvest did not show obvious rhizomania symptoms but

TEST B799. HYBRID PERFORMANCE OF MONOGERM S₁ PROGENY LINES UNDER RHIZOMANIA, IMPERIAL VALLEY, CA., 1998-99

72 entries x 1-row plots,	4 reps., RCB, 6 blocks per rep 18 ft. long, 24 blocks, 12 rows					Planted: Harvested:	September May 12,	ber 23-24 12, 1999	, 1998
Variety	Description	Acre	Yield Beets	Sucrose	Beets/ 100'	Bolters	Root Rot1	Clean Beets	NO3-N
		sqT	Tons	æ	No.	∞ I	o⁄0 	∞ 1	Mean
Checks	Spreckels, 9-98, L1162401	8420	32.17	0	152	2.7	0.0	88.4	
X869H37	\sim	8717	3.4	3.0	158			6	
B4776R	Beta 4776R.7653 (3-27-98)	7967			132	2.2	0.0	89.3	45
X869H50	C790-15CMS x Y769	6872	1.0	1.0	149	•		9	
S, lines from	1 popn-833								
		8130	33.00	ო.	153		0.0	•	42
X869H5	5833-5aa (C833-5)	\vdash	9	1.8	140	•		7	25
Y869H33-1	7833-1aa x Y769	6453	30.77	10.52	152	1.9	0.0	90.6	18
X869H33-3	7833-3aa x Y769	0	34.38	1.6	136	•	•	0	27
X869H33-10	7833-10aa x Y769	7162	8.4	2.6	5	•	•	9.	26
Y869H33-11	7833-11aa x Y769	7193	0	1.8	Э	6.0		Η.	23
Ү 869Н33-12	7833-12aa x Y769	6564	۲.	11.68	136	0.0	5.8	93.9	27
Ү869 H12	5833-12aa (C833-12) x Y769	7298	0	1.9	ന	0.0	•	Η.	12
S_1 lines from	. popn-834								
X869H29	5829-3aa (C829-3) x Y769	6340	5.3	2.4	147	•	0.0	Э.	30
X869H34-1	7834-1aa x Y769	6650	29.97	11.24	149	0.0	0.0	90.7	35
X869H34-2	7834-2aa x Y769	7297	0.7	1.8	140	•		Ή.	19
Y869H34-3	7834-3aa x Y769	6658	8.3	1.7	4	•	0.0	ω.	39
Y869H34-5	×	6625	29.42	11.22	158	2.5	0.0		28
X869H34-8	7834-8aa x Y769	7972	3.7			•	0.0	90.6	22
S1 lines from	. popn-829								
Y869H28-9	7828-9aa x Y769	7673	32.30	11.85	147	0.0	0.0	88.3	18
Y869H28-10	7828-10aa x Y769	7742	1.4	ю.	143	0.0	•	7.	15

TEST B799. HYBRID PERFORMANCE OF MONOGERM S₁ PROGENY LINES UNDER RHIZOMANIA, IMPERIAL VALLEY, CA., 1998-99

(cont.)

		Acre	Acre Yield		Beets/		Root	Clean	
Variety	Description	Sugar	Beets	Sucrose	1001	Bolters	Rot	Beets	NO3-N
		sqT	Tons	%	No.	ογ∘I	%	œ۱	Mean
,,	ω.		,	,					!
х869Н69	7869aa x Y769	6469	8.6	1.3	9	•	٠		25
Y869H69- 1	7869- 1aa x Y769	8299	6.5	1.3	4	•	•	92.6	27
X869H69- 2	7869- 2aa x Y769	6926	28.76	12.06	147	1.6	0.0	92.4	35
X869H69- 4	7869- 4aa x Y769	7032	8.9	2.1	സ	•	0.0	84.8	20
Х869Н69- 5	7869- 5aa x Y769	7300	9.0	1.9	138	•	•	4.	25
7869H69- 6	7869- 6aa x Y769	7543	9.0	2.3	5	•	•	ö	46
7 -69H698Y	7869- 7aa x Y769	6730	29.16	11.54	139	0.0	1.0	0.06	22
х869н69-13	7869-13aa x Y769	7348	1.8	1.5	4	•	•	1.	35
01 000000	7	0	7		Ú		c	c	cc
ET-EOUEOOI	×	6001			Ó	•	•	,	77
X869H69-20	7869-20aa x Y769	8069	8.5	2.1	Ŋ	•	٠		21
X869H69-20B	7869-20Baa x Y769	0689	30.92	11.16	135	0.0	0.0	89.7	34
Y869H69-24	7869-24aa x Y769	7725	1.1	2.4	3	0.0	•	2	20
S_1 lines from	936-ngod								
	7838aa x Y769	10	2.8	2.3	3	•	•	<u>ი</u>	40
X869H36- 3	7836- 3aa x Y769	7270	30.57	11.88	115	0.0	0.0	89.5	36
Y869H36-11	7836-11aa x Y769	78	2.9	1.9	3	•		ი	17
X869H36-14	7836-14aa x Y769	7313	0.2	2.0	0	•	0.0	:	23
S, lines from	popn-837 85.9								
Y869H77-1		7048	8.6	2.3	2	•	•	δ.	
Y869H77-1B	7837-1Baa x Y769	6756	28.60	11.83	126	0.0	0.0	87.5	29
X869H77-2	7837-2aa x Y769	6750	6.4	2.7	4	•	•	5.	
X869H77-3	7837-3aa x Y767	7004	7.8	2.6	Ω	•	•	ω.	
X869H77-4	7837-4aa x Y769	5591	4.4	1.4	7	•	•	7 .	

TEST B799. HYBRID PERFORMANCE OF MONOGERM S₁ PROGENY LINES UNDER RHIZOMANIA, IMPERIAL VALLEY, CA., 1998-99

(cont.)

Variety	Description	Acre	Acre Yield ar Beets	Sucrose	Beets/ 100'	Bolters	Root Rot1	Clean Beets	NO3-N
		Ips	Tons	%	No.	o∻	% I	% 1	Mean
S ₁ lines from popn-839	1 popn-839								
Y869H79-1	$7839-1aa \times Y769$	O.	30.35	ij	147				28
Y869H79-2	7839-2aa x Y769	ø	2.3	1.6	Ţ	0.0	•	8	18
X869H79-3	7839-3aa x Y769	6736	2.0		139	0.0	0.0	92.0	23
X869H79-4	7839-4aa x Y769	6706	9.1	1.5	4	0.0	•	о О	36
X869H79-5	7839-5aa x Y769	7935	34.14	11.62	143	0.0		ά.	35
X869H79-5B	7839-5Baa x Y769	σ		ij	135	0.0	2.3	92.7	13
X869H79-6	7839-6aa x Y769	4	26.62	11.97	133	0.0		ė.	24
X869H79-10	7839-10aa x Y769	7492	1.8	ij	136	•	•	ъ.	30
S ₁ lines from	popn-831-4								
Y869H4	5831-3aa (C831-3) x Y769	7362	0.8	1.9	122	•		2	28
Y869H27-1	$7831-4-1aa \times Y769$	7200	31.78	11.29	124	0.0	0.0	89.4	26
Y869H27-2	$7831-4-2aa \times Y769$	7506	3.7	1.1	128	•	•	9.	46
Y869H27-7	7831-4-7aa x Y769	6465	9.0	11.15	133		•	ص	30
X869H27-8	$7831-4-8aa \times Y769$	8824	7.2	σ.	131	•	•	2	30
Y869H27-9	7831-4-9aa x Y769	6077	26.96	11.31	118	0.0	0.0	91.1	19
Y869H27-10	$7831 - 4 - 10aa \times Y769$	7373	2.6	1.3	125	•	•	7	40
S ₁ lines from	808-udod								
Y869H9-1	7808-1aa x Y769	7734	1.7		136	0.0		•	21
х869н9-2	7808-2aa x Y769	7298	•	12.23	135	0.0	0.0	92.7	45
X869H9-3	7808-3aa x Y769	0	9.1	ω.	126	1.1	•	•	31
Ү 869н9-4	7808-4aa x Y769	7142	7.5	3.1	138	•	0.0	ω	29
X869H9-7	7808-7aa x Y769	0	7.7	1.3		•		ω.	31
х869н9-8	7808-8aa x Y769	6337	29.46	10.78	138	0.0	0.0	92.2	36
X869H9-9	7808-9aa x Y769	6415	5.2	2.7	cr)	•	•	9	50

HYBRID PERFORMANCE OF MONOGERM S₁ PROGENY LINES UNDER RHIZOMANIA, IMPERIAL VALLEY, CA., 1998-99 TEST B799.

(cont.)

		Acre Yield	ield		Beets/		Root	Clean	
Variety	Description	Sugar	Beets	Sucrose	1001	Bolters	Rot^1	Beets	NO3-N
		sqT	Tons	% 1	No.	%	%	%	Mean
S ₁ lines from	S ₁ lines from popn-808 (cont.)								
Y869H9-12	7808-12aa x Y769	8216	33.96	12.07	150	0.0	0.0	88.8	32
Y869H9-13	7808-13aa x Y769	6974	29.32	11.77	146	0.0	1.9	91.0	48
Ү869Н9-16	7808-16aa x Y769	5174	23.32	11.02	133	0.0	1.0	9.68	12
S ₁ lines from popn-818	popn-818								
Y869H15-1B	6818-1Baa x Y769	7908	32.96	12.01	143	0.0	0.0	89.5	30
Y869H15-2B	6818-2Baa x Y769	7064	30.71	11.48	147	0.0	2.1	88.3	12
Y869H15-1	6818-1aa x Y769	6631	26.02	12.65	146	0.0	8.3	90.5	21
X869H15-2	6818-2aa x Y769	8438	34.47	12.23	147	0.0	0.0	86.1	13
X869H15-6	6818-6aa x Y769	7618	30.67	12.44	136	0.0	3.1	90.4	23
X869H15-21	6818-21aa x Y769	6928	27.79	12.48	142	0.0	0.0	88.1	33
Mean		7183.2	30.3	11.86	140.0	0.2	0.4	89.5	27.6
LSD (.05)		1342.3	5.1	96.0	22.7	1.6	2.2	4.5	29.7
C.V. (%)		13.4	12.1	5.79	11.6	507.8	401.6	3.6	77.0
F value		2.1**	2.2**	2.68**	1.9**	1.3NS	2.7**	2.2**	0.8NS

NOTES: See Test B699. Test B799used as screen to identify monogerm lines with adaptation under rhizomania conditions to Imperial Valley.

¹Root rot. See Test B499.

TEST B899. HYBRID PERFORMANCE OF MULTIGERM PROGENY LINES UNDER RHIZOMANIA, IMPERIAL VALLEY, CA., 1998-99

1998	NO3-N	Mean	115	150	88	109		123	117	136	118	122	113	125	113	157	136		120	107	111	112
September 24, May 11, 1999	Clean Beets	o/e	6	89.8 89.4	•	90.3			88.0		ω.	÷.	0.68	7.	رى	89.1	6		88.8	87.4	•	
Septe d: May	Root Rot ¹	o⁄o I		o o o		0.0			0.0		•	•	0.0	•	•	0.0	•		0.0	0.0		
Planted: Harvested:	Bolters	o,0		6.0 0.0	•	0.0		0.0	•	0.0	0.0		0.0	•	•	0.0	•		0.0	0.0	•	•
	Beets/ 100'	No.	147	157 154	150	153		153	131	149	142	139	150	136	139	150	150		142	157	146	152
	Sucrose	00 	2.7	11.93 13.22	3.3	13.99		12.94	•	12.41	ĸ.	. 7	12.32	2.1	•	12.67	•		0.	12.55	3.5	2.7
	Yield Beets	Tons	م	32.82 33.26	Ή.	32.98		ø.	28.33	о О	о О	o.	33.62	o.		29.74	ė.		4.	37.35	ഹ	σ.
	Acre Sugar	I.bs	7593	7787 8820	8270	9240		9969	6229	7291	7595	7130	8290	7883	7813	7458	8394		7363	9372	8547	7150
72 entries x 4 reps., RCB, 6 blocks per rep 1-row plots, 18 ft. long, 24 blocks, 12 rows	Description		Betaseed, 7-10-97	Spreckels, 9-98, L782402 Holly HH108, 9-3-97	Spreckels, 9-98, L1162401	Betaseed 4776R.7653 (3-27-98)	N Bvm, R22, C51 resistance	C790-15CMS x RZM R735 (C79-7)	C790-15CMS x RZM R736, R746 (C79-8)	C790-15CMS x RZM R754	C790-15CMS x RZM Y773	C790-15CMS x RZM R779 (C79-1)	C790-15CMS x RZM Y767 (C67)	C790-15CMS x RZM Y771	C790-15CMS x RZM-% Y672	C790-15CMS x RZM Y775,	6831-4HO x RZM Y775,	Dr MM of De lines		x 6925-19	C790-15CMS x RZM-ER-% 6913-7	C790-15CMS x RZM-ER-% 6911-4-10
72 entries x 1-row plots,	Varietv		Checks B4035R	SS-778R Rizor	Rifle	B4776R	Hybrids with	R835H50	R836H50	R854H50	Y873BH50	R879H50	X867H50	Y871H50	X872H50	X875H50	Y875H27	אלינה מה ייילינו לדינה מה ייילינו	8931H50	8925-19H50	8913-70H50	8911-4-10H50

(cont.)

Varietv	Description	Acre	Yield Beets	Sucrose	Beets/	Bolters	Root Rot ¹	Clean Beets	NO3-N
		Tps	Tons	o⊱l	No.	₩	%	ઝ ∘ I	Mean
Hybrids with	Rz,MM,S ^f ,Aa lines (cont.)	00	7.0	2.5	129	0.0	0.0	6	77
8918-21H50		6609	23.31	13.05	125		•	86.3	120
Z825-6H50	C790-15CMS x Z625-6	0	4.0	2.8	152	1.6	0.0	ω.	134
Z825-9H50	×	3	6.3	3.9	ϵ	•	•	7.	105
Z830-11H50	C790-15CMS x Z630-11	8321	34.44	12.10	145	0.0	0.0	85.4	102
CR812H50	C790-15CMS x RZM CR712	7163	29.71	12.13		6.0	0.0	88.8	127
CR813H50	C790-15CMS x RZM CR713	8647	5.3	2.2	164	2.4	•	5	124
Hybrids with	Hybrids with R22(C51),MM,S ^f ,Aa lines			,	,				,
8926н50	C790-15CMS x RZM 7926	7927	29.86	13.24	152	ი.0	0.0	85.7	68
8926H55	7835H50 x RZM 7926	4	9.8	2.5	150	1.9	٠	0	133
8926H58	7838H50 x RZM 7926	8793	5.5	2.3	167		0.0	<u>.</u>	115
8927-29H50	C790-15CMS x 6927-29	m	0.5	2.9	140	•	•	œ.	107
8927-30H50	C790-15CMS x 6927-30	Н	35.07	13.05	153	0.0	0.0	86.0	93
8927-33H50	C790-15CMS x 6927-33	\mathbf{S}	9.9	3.1	150	•	•	7 .	112
8927-37H50	C790-15CMS \times 6927-37	75	2.0	2.1	2	•	0.0	о О	104
Hybrids with R878H50	C790-15CMS × R778.R778%	47	2.3	3.0	140		0.0	ω	104
8930- 19H50	×	8569			142	0.0		86.3	110
8930- 39H50	C790-15CMS x 6930- 39	75	3.9	2.8	143	•		œ	101
8930-102H50	$C790-15CMS \times 6930-102$	57	9.3	2.8	138	•	•	4.	122

TEST B899. HYBRID PERFORMANCE OF MULTIGERM PROGENY LINES UNDER RHIZOMANIA, IMPERIAL VALLEY, CA., 1998-99

		Acre	Yield	000000000000000000000000000000000000000	Beets/	+ C + C +	Root Pot1	Clean	N CON
Aditec	pesci ipcioni	I.bs	Tons	% 	No.	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	o ∞	2 0 ∞	Mean
Hybrids with	C76,Rz,MM, S ^f ,Aa lines								
R876-89-5H50	C790-15CMS x RZM-% R576-89-5	8663	2.2	3.5	154	0.0	0.0	87.2	91
R882H50	C790-15CMS x R781,R776	33		12.58	131	0.0	0.0	•	151
R882H27	6831-4HO x R781,R776	8282		2.3	135	2.0	0.0	91.8	105
8929- 41H50	C790-15CMS x 6929- 41	25	4.	3.5	5	•		7.	\vdash
8929- 72H50	C790-15CMS x 6929- 72	84	3.2	3.2	153	•	•	0	119
8929-102H50	C790-15CMS x 6929-102	8619	34.68	12.39	147	0.0	0.0	93.0	66
8929-112H50	$C790-15CMS \times 6929-112$	30	4.6	3.4	147	•	•	0.	95
8929-114H50	×	95	0.7	4.5	ന	0.0	•	6.	106
8929-115H50	×	66	8.7	2.9	4	•	0.0	0	94
8929-133H50	×	03	5.4	3.8	4	0.0	•	6	ч
929	$C790-15CMS \times 6929-153$	8797	33.41	13.19	156	0.0	0.0	89.5	108
8929-154H50	C790-15CMS x 6929-154	51	8.1	1.6	2	0.0	0.0	8	4
Monogerm lines	s topcrossed to C69								
X869H50	Ms x	7508	9.5	2.6	4	•	•	ij	$^{\circ}$
X869H27	6831-4HO x Y769	7618	31.07	12.24	152	0.0	6.0	90.3	125
X869H35	7835mmaa x Y769	91	8.3	2.1	4	•	•	2	2
х869нзв	7838mmaa x Y769	7624	9.0	3.0	143	0.0	0.0	•	⊣
X869H17	7817HO x Y769	7754	0.8	2.5	r	0.0	0.0	ί.	112
Y869H9- 1	7808- 1aa x Y769	6833	29.04	11.81	154	0.0	0.0	91.3	86
¥869H9- 2	7808- 2aa x Y769	6011	4.2	2.4	142	0.0	0.0	6	106
хв69н9- з	7808- 3aa x Y769	37	8.0	3.1	4	0.0	•		06
Y869H9- 4	7808- 4aa x Y769	7273	27.32	13.30	157	0.0	0.0	6.68	88
х 1869но- 7	7808- 7aa x Y769	4	6.7	2.6	4	0.0	•	ij.	109
8 -6H698Y	7808- 8aa x Y769	5756	26.52	10.85	139	c	c	7 10	132
	:)	•	•)

TEST B899. HYBRID PERFORMANCE OF MULTIGERM PROGENY LINES UNDER RHIZOMANIA, IMPERIAL VALLEY, CA., 1998-99

(cont.)

		Acre Yield	rield		Beets/		Root	Clean	
Variety	Description	Sugar	Beets	Sucrose	1001	Bolters	Rot^1	Beets	NO3-N
		sqT	Tons	o⊱1	No.	%	o⊳1	o o	Mean
Monogerm lines	Monogerm lines topcrossed to C69 (cont.)								
Х869Н9- 9	7808- 9aa x Y769	6336	25.10	12.47	142	0.0	0.0	9.68	102
Y869H9-12	7808-12aa x Y769	7307	30.17	12.07	150	0.0	0.0	88.9	115
Y869H9-13	7808-13aa x Y769	6386	25.10	12.78	142	0.0	0.0	93.8	130
х 869н9-16	7808-16aa x Y769	5858	25.01	11.81	120	0.0	0.0	91.9	118
Y869H15-1B	6818-1Baa x Y769	7186	27.35	13.08	157	0.0	0.0	89.7	127
Y869H15-2B	6818-2Baa x Y769	6489	28.66	11.85	152	0.0	0.0	7.06	118
Y869H15-1	6818-1aa x Y769	6357	24.79	12.76	142	0.0	6.0	92.2	105
Y869H15-2	6818-2aa x Y769	7314	29.16	12.53	152	0.0	0.0	86.9	103
Y869H15-6	6818-6aa x Y769	6782	\vdash	12.47	129	0.0	0.0	88.2	113
Y869H15-21	6818-21aa x Y769	5730	22.93	12.60	132	0.0	2.8	89.1	114
Y869H18	7818HO x Y769	7106	29.07	12.19	160	0.0	0.0	88.3	129
Y869H49	7848H88mmaa x Y769	7162	28.48	12.56	156	0.0	0.0	90.4	116
Mean		7705.6	30.36		146.0	0.4	0.1	88.8	114.1
ISD (.05)		1574.8	5.47	1.16	22.5	2.4	1.5	4.1	42.7
C.V. (%)		14.7	12.92	6.56	11.1	394.9	858.1	3.3	26.9
F value		3.0**	* 3.05**	2.20**	1.3NS	1.5*	1.4NS	3.4**	1.0NS

NOTES: See Test B699. Test B899 used as screen to identify multigerm lines and progeny with adaptation under rhizomania conditions to Imperial Valley.

¹Root rot. See Test B499.

TEST B599. AREA 5 CODED RHIZOMANIA YIELD TEST, IMPERIAL VALLEY, CA., 1998-99

8 reps, RCB(E) 27 ft. long Variety Source	Source		Acre Sugar	Yield Beets Tons	Sucrose	Planted: Harvested Beets/ 100'	October 21: June 10, Clean Beets	, 1998 1999 NO3-N Mean
7067321		Betaseed	7845	ω.	9. 8.	167		210
Rizor		Spreckels	7590	25.73	14.74 14 52	167	94.3 o	193
JCG7322		Betaseed	7485		3.4	168		228
8CG7064		Betaseed	7052	51	13.90	168	4.	195
Beta 4035R		Betaseed	6119	2.	w.	167	•	201
SS-NB7R		Spreckels	6227	23.79	13.08	159	•	200
Beta 4776R		Betaseed	5997	÷.	3.7	159	95.2	276
Rifle		Spreckels	5856	19.62	14.88	167	•	203
Alpine		Spreckels	6151	•	•	160	94.2	205
SS-778R		Spreckels	5850	22.58	ω.	164	•	240
Beta 4684R		Betaseed	5757	•	7.	162	94.4	214
Phoenix		Spreckels	5679	÷.	12.96	168	94.9	320
97CX01		Spreckels	5630	i.	•	166	m	196
Summit		Spreckels	5741	22.27	12.97	154	93.4	212
7KJ0191		Betaseed	5493	o.	14.28	163	\vdash	228
SS-781R		Spreckels	5621	ά.	•	162	94.5	240
98HX853		Spreckels	5508	20.93	13.17	160	94.6	218
Beta 4210R		Betaseed	5877	25.46	. 4	162	95.6	374
Pinnacle		Spreckels	5429	20.91	12.99	159	94.5	245
5CG7540		Betaseed	5329	21.39	٠. ت	165		317
Rival		Spreckels	4686	17.21	13.64	156	92.8	196
Imperial		Spreckels	4855	œ.	12.78	154	94.6	250

TEST B599. AREA 5 CODED RHIZOMANIA YIELD TEST, IMPERIAL VALLEY, CA., 1998-99

(cont.)

			Acre Yield	ield	,	Beets/	Clean	
Code	Variety	Source	Sugar	Beets	Sucrose	No.	Beets 1%	Mean Mean
98IVR -14 - 8	Beta 4684 US H11	Check Standard	4625 4392	16.61 17.64	13.97	160 133	93.1 92.4	201
- 6 -13	7CG7400 SS-IV2	Betaseed Check	4146 3993	15.55 16.37	13.06 12.25	162 167	90.2 92.4	271 204
USDA entries	C790-15CMS x RZM 7926	7926	665 655	25.10	13.21	156	93.2	800
Y875H50	C790-15CMS x RZM Y775	Y775	6012	22.67	13.25	164	94.1	224
Y875H27 8926H37	C831-4HO x RZM Y775 4807HO (C306/2CMS) x R	75 3) × RZM 7926	5748 5607	22.13 24.38	13.03 11.52	153 153	93.2 93.4	190 249
Y875H37	4807HO (C306/2CMS) x R) x RZM Y775	5525	22.41	12.24	160	93.2	246
Mean			5808.9	21.88	13.24	161.1	93.6	233.0
LSD (.05) C.V. (%)			1114.8 19.5	11.89 18.89	0.61 4.68	11.4 7.2	1.6 1.8	46.2 20.1
F value			5.6**	* 4.76**	14.64**	3.0**	4.0**	6.5**

TEST B599. AREA 5 CODED RHIZOMANIA YIELD TEST, IMPERIAL VALLEY, CA., 1998-99

(cont.)

		Recover.	Recover.	Recover.	Known	;		;	
Code	Variety	Sugar	Sugar	Sugar	Sugarloss	Sodium	Potassium	NH2-N	Impur.
		1bs/a	1bs/t	o(0	<u>lbs/a</u>	wdd	wdd	wdd	Value
98IVR -21	7CG7321	6578	232	83.6	1267	766	2493	566	15103
-30	Rizor	6551	254	ė.	1039	695	2339	568	13673
en I	Beta 4430R	6393	253	87.2	950	837	2133	439	12432
-17	7CG7322	6311	227	4	1174	881	2395	512	13936
-29	8CG7064	6229	244	87.8	823	707	1905	426	11283
-22	Beta 4035R	5248	235		931	006	37	519	14018
-18	SS-NB7R	5187	217	83.0	1039	908	53	597	82
- 2	Beta 4776R	5099	235	85.0	868	962	2147	N	13715
-28	Rifle	5097	259	87.1	759	657	22	523	12827
-26	Alpine	4992	207	•	1159	768	2612	957	18307
-27	SS-778R	4957	216	4.	892	899	2369	457	13405
-24	Beta 4684R	4939	247	85.5	817	954	2250	530	13997
-16	Phoenix	4825	219	4.	854	905	2361	469	13527
-11	97CX01	4816	223	85.4	813	825	2112	457	12505
- 7	Summit	4811	217	е Э	930	808	2577	497	13998
-25	7KJ0191	4719	243	84.8	774	875	2439	523	14128
თ 1	SS-781	4686	209	82.7	935	913	2372	544	14295
i N	98HX853	4683	224	84.9	824	798	2257	510	13282
-12	Beta 4210R	4666	182	•	1211	1150	57	571	15890
-10	Pinnacle	4578	219	84.1	851	1056	2259	461	13723
7 -	5CG7540	4369	206	81.8	096	1166	ന	535	14922
-20	Rival	4105	239	87.6	582	725	2075	382	11350
-19	Imperial	4079	214	83.6	776	1025	$^{\circ}$	504	13971

(cont.)

Vari	Varietv	Recover. Sugar	Recover. Sugar	Recover.	Known	Sodium	Potassium	ארי או	Tmciir
		<u>lbs/a</u>	1bs/t	₩	lbs/a	wdd	wdd	i	Value
ta	Beta 4684	3944	238	85.1	680	876	2197	554	13820
US H11	н	3734	209	84.4	629	784	2134	493	12765
7CG7400	00	3468	217	83.1	678	1016	2354	554	14706
ss-IV2	2	3230	199	81.2	762	1086	2462	548	15163
		5633	223	84.3	1023	748	2471	514	13676
		5059	223	84.0	953	865	2314	557	14105
		4880	221	84.6	869	775	2410	471	13209
		4862	185	80.3	1172	983	2945	537	15910
		4506	199	80.9	1019	1021	2637	551	15400
		4913.6	222.9	84.0	908.6	889.5	2352.3	526.5	13995.8
		1023.2	16.7	3.7	238.1	281.3	417.9	228.5	3014.8
		21.1	7.6	4.5	26.6	32.1	18.0	44.1	21.9
		5.2**	**6.6	2.7**	3.0**	1.7NS	1.8NS	1.3NS	1.6NS

collapsing and test was not uniform in appearance. It appeared that rhizomania and other soil-borne problems were NOTES: Test was under high nitrogen status. Due to initial emergence and stand problems, test was replanted on October 21, 1998. Up to mid-May, test appeared uniform. At harvest, some plots (e.g., entries 8,13 & 14) were moderate but variable across the field. USDA entries Y775 and 7926 segregate for resistance to rhizomania from Beta vulgaris ssp. maritima thru C50 (R22).

EVALUATION OF TESTCROSS HYBRIDS FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, CA., 1998-99 TEST B1099.

			(
Variety	Description	Stand Count	% Bolting	App	Appearance S	Score	Living Plants
		No.	06/11	05/13	06/11	07/08	o,⇔
Checks	100-10-61 6321 93611 800000	_	~				d
D4 / / OK	2		* C	•			
Diflo	90-0 SIG45		0.0				. α
R522 (Sp)	, , , , (C	20.3	14.1	1.8	1.5	1.0	73.1
Topcrossed to	o C69						
Х869Н38	7838aa x Y769	o.	0.0	4.0	•	4.5	9
X869H27	6831-4HO x Y769	ω.	•	•	•		4.
Y869H19	7818H50 x Y769	18.8	0.0	3.5	3.3	3.8	27.9
X869H18	7818HO x Y769	o.	•	•	•	•	ė.
X869H20	7818-4H50 x Y769	7.	0.0	•		4.5	o.
Y869H21	7818-14H50 x Y769	20.3	0.0	3.3	3.8	4.0	18.7
X869H22	7818-22H50 x Y769	80	0.0	•	•	4.3	7.
х869H23	7818-23H50 x Y769	o,		•	•	4.3	9.
X869H15-1B	6818-1Baa x Y769	•	0.0	•	•	4.3	16.4
X869H15-2B	6818-2Baa x Y769	18.3	0.0	3.3	3.8	4.5	10.6
X869H15-1	6818-1aa x Y769	•	0.0	•	•	4.8	•
Y869H15-2	6818-2aa x Y769	5.	•	•	•	4.0	17.7
X869H15-6	6818-6aa x Y769	9.		4.0	•	4.8	
X869H15-21	6818-21aa x Y769	8	0.0	•	•	•	6.3
X869H9-1	7808-1aa x Y769	19.5	0.0	3.5	3.5	4.3	13.0
х869н9-2	7808-2aa x Y769	o.	0.0	•	•	4.8	•
х869н9-3	7808-3aa x Y769	ω.		•	•	•	9
X869H9-4	×	18.5	0.0	3.8	3.5	4.0	29.5
X869H9-7	7808-7aa x Y769	9.	•	•	•	•	9
Y869H9-8	0377 *0-0007	1					c

EVALUATION OF TESTCROSS HYBRIDS FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, CA., 1998-99 TEST B1099.

(cont.)

7 7 7 4 4		Stand	oko -	•		,	Living
Variety	Description	No.	06/11	Appe 05/13	Appearance score (13 06/11 07/0	07/08	Plants
Topcrossed to C69 (cont.)	C69 (cont.)						
6-6Н698Х	7808-9aa x Y769	16.5	0.0	3.5	3.5	3.8	28.3
X869H9-12	7808-12aa x Y769	20.3	0.0	3.3	3.3	4.0	30.8
X869H9-13	7808-13aa x Y769	18.0	0.0	4.0	3.3	3.8	22.6
Х869Н9-16	7808-16aa x Y769	14.3	0.0	4.0	3.8	4.3	10.9
Testcrosses to	Testcrosses to Y75 and popn-926						
Y875H27	_	17.5	0.0	2.5	3.0	3.8	33.8
X875H50	C790-15CMS x RZM Y775,	18.5	0.0	2.3	2.3	2.5	52.3
8926H50	C790-15CMS x RZM 7926	19.8	0.0	2.3	2.3	3.0	39.3
X872H50	C790-15CMS x RZM-% Y872	18.3	0.0	1.8	1.8	2.3	77.2
Mean		18.3	0.5	3.4	3.3	3.9	26.2
LSD (.05)		3.0	3.4	8.0	0.7	0.8	18.3
C.V. (%)		11.8	437.8	16.7	15.8	13.9	49.8
F value		3.3**	4.5**	5.3**	5.8**	9.5**	8.7**

NOTES:

Appearance scored on a scale of 1 to 5 where: 1 = very good canopy; 2 = good canopy and appearance often segregating; 3 = intermediate and variable; 4 = fair; and 5 = poor to mostly dead plants.

However, other factors such as plant vigor, cyst nematode infection, root rots, etc. could have influenced vigor, number of dead leaves, and dead plants. The assumption was that plant health and appearance was Appearance scored relative to the overall test at time and based upon canopy size, uniformity, color, mostly being influenced by reaction to rhizomania and rhizomania under high temperature conditions. appearance.

Coefficients of correlation for % Living plants vs. Appearance scores for 5/13, 6/11, & 7/8 and Stand Counts (October 98) are r = -.55**, -.75**, -.90**, and .21*, respectively. Stand counts made post thinning in October 98 and living plants counted 08 July 1999.

EVALUATION OF MULTIGERM LINES FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99 TEST B1199.

Planted: September 24, 1998 Living Plants 20.5 37.3 42.5 38.3 39.4 13.4 33.8 37.8 41.7 12.1 38.8 11.5 71.3 67.4 46.7 Not harvested for yield 80/10 4.3 1.0 2.5 8.2 3.0 3.0 3.5 $\overset{\circ}{\omega}$ $\overset{\circ}{\omega}$ $\overset{\circ}{\omega}$ 3.5 3.5 4.3 3.3 3.8 Appearance Score 06/11 3.0 1.3 2.5 3.0 3.0 3.3 3.0 3.0 3.9 3.3 2.8 2.8 3.3 4.0 05/13 4.0 1.8 3.5 4.0 3.0 3.0 3.0 3.0 Bolting 06/11 0.0 0.0 0.0 2.3 0.00 0.0 20.9 0.0 0.0 1.5 0.0 19.3 Stand 15.3 18.3 19.3 19.3 19.5 22.0 18.8 18.3 18.3 20.3 16.0 18.3 20.3 19.3 18.8 19.0 20.5 19.8 17.3 Count N S (C79-3)Betaseed 4776R.7653 (3-27-98) RZM-% R576-89-5NB, (C76-89-5) Spreckels, 9-98, L1162401 Spreckels, 9-98, X782402 64 entries x 4 replications, sequential (C79-2), R725 (C82) Inc. R781,R776,... (C82) $(C80Rz \times SR)$ (C80Rz \times SR) Description RZM Y775,Y773,Y772,... (C79-7, SES) RZM-%S R322R4,...(C51) но11у ни108, 9-3-97 R776, R781,... RZM R778% (C78) R780 (C80) RZM Y769 (C69) 1-row plots, 13 1/2 ft. long EL#S RZM EL#s R724 **RZM Y775** RZM Y768 R735 RZM 7926 1997 RZM RZM RZM RZM RZM MM, O.P. lines R876-89-5NB Variety Y875 (Iso) (osI) 698X 3926 (Iso) R522 (Sp) Y875 (Sp) 98-EL-02 98-EL-04 SS-778R Checks B4776R US H11 Rifle Rizor R878% R880 **X868** R824 R835 R882 R881

TEST B1199. EVALUATION OF MULTIGERM LINES FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99

Variety Description Count Bolting Appearance Score Plan No. $06/11$ $05/13$ $06/11$ $07/08$ $\frac{8}{2}$							
Count Bolting Appearance Score No. 06/11 05/13 06/11 07/08			Stand	ф			אלו
Count Bolting Appearance Score No. 06/11 05/13 06/11 07/08							
No. 06/11 05/13 06/11 07/08			1 11110	11:11		41000	ב
06/11 05/13 06/11	variety	Describinon	Conne	POTEING	Appearance	e acore	ช - น
			ON	06/11	05/13 06/1	1 07/08	o)¢
							1

Variety	Description	Stand	% Bolting	Appe	Appearance Score	Score	Living Plants
		No.	06/11	05/13	06/11	07/08	o+>
MM, O.P. lines (cont.)	(cont.)						
97-C37	Inc. U86-37	17.5	0.0	4.3	4.0	4.5	7.2
R879	RZM R779 (C79-1, RZ)	17.3	0.0	4.8	4.5	4.5	6.8
R836	RZM R736 (C79-8, R22)	21.3	6.2	2.3	2.5	2.8	29.0
US H11	1997	15.8	0.0	4.3	•	4.3	14.2
R746 (Iso)	RZM R646	18.0	0.0	3.3	9. 3	3.8	26.4
R853	RZM-ER-% R653	20.3	0.0	4.0	3.5	4.3	0.6
R854	RZM R754	19.0	0.0	3.8	3.8		9.5
Y873	RZM-ER-% Y673	17.8	0.0	2.0	•	3.0	39.8
Y873B	RZM Y773	20.0	0.0	2.5	2.8	3.5	32.3
97-C37	Inc. U86-37	20.3	0.0	4.8	•	4.5	6.3
R840	RZM R740 (C79-#s)	21.5	1.1	2.3	3.0	•	32.7
X866	RZM Y766	20.3	1.1	1.8	1.5	2.0	42.4
X867	RZM Y767 (C67)	0	0.0	•	2.3		ი
Y871	RZM Y771	19.8	1.0	2.5	2.0	2.8	34.5
Y872	RZM-8 Y672	19.8	0.0	•	2.5		m.
Y872B	RZM Y772 (C72)	19.8	1.1	2.8	2.8		29.7
Y875 (Iso)	RZM Y775	œ	1.5	•			32.2
8810M	RZM 7810NB (C890-#s)	•	0.0	3.8	3.3		m.
R826	RZM R726 (C26)		10.1	2.5	2.5	3.3	25.7
R827	RZMR727A,B (C27)	20.5	1.1	3.0	3.0		27.3

TEST B1199. EVALUATION OF MULTIGERM LINES FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99

(cont.)

Varietv	Description	Stand Count	% Bolting	Арре	Appearance 8	Score	Living Plants
		No.	06/11	05/13	06/11	07/08	o∞
MM, O.P. lines	lines (cont.)						
P811	RZM-PMR 6203,6208(C),(R79 \times P03,P04)	18.8	10.9	3.0			38.3
	6211,6217(C), (C918 x	6		1.0		•	
	03)	19.0	0.0	4.8	4.3	4.8	1.3
	6205,6206(C),(C37 x	9	0.0	4.0	•	4.5	•
MM, Sf, Aa popul	populations						
	Inc. N623, N624 (galls), SBCN resist.	7	0.0	3.0			18.5
CR811	RZM CR711 (CR09/10)	18.3	•	3.0		3.8	25.7
CR812	RZM CR712 (931 x CR11)	\sim	•	3.0	3.3	3.8	27.7
CR813	RZM CR713 (932CT x CR11)	17.3	0.0	3.0		4.3	20.8
7777	The FOAT (2001)	7	c	~	-	α	ر. 1
	, ,	r C	•	•	•) L	
		O 0) i	n 0	o (# (C	# · ·
	RZM 7926aa x A	`	•	٠	٠	7.8	39.4
8926 (Sp)	7931aa x RZM 7926	0	0.0	•	2.8	2.8	36.1
8927-29	, Inc.	Ŋ	0.0	5.0	•		0.0
	, (5921H18), Inc.		0.0	•	•	•	。
8927-33	6927-33 (A, aa)	18.8	0.0	2.5	2.5	3.0	32.3
	, (5921H18), Inc.		0.0	•	•	•	•
8924	RZM 7924aa x A	o.	•	э. Э.		4.0	
	7932CT,7201-7215aa \times A, R_z -CTR	15.5	0.0	3.5	4.3	4.8	3.6
2831	Ø	7.	•	3.0	3.8	4.3	12.5
8935 (Iso)	RZM R776-89-5H13	17.8	0.0	3.8	3.8	4.3	11.1

EVALUATION OF MULTIGERM LINES FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99 TEST B1199.

, to	, + + · · · · · · · · · · · · · · · · ·	Stand	0 % † ₹	6		, ,	Living
Variety	Describution	No.	06/11	05/13	5/13 06/11 07/0	07/08	8
MM, St, Aa por	MM,Sf, Aa populations (cont.)						
8936	RZM R776-89-5H31	19.3	0.0	3.3	3.0	4.8	4.7
8937	RZM R776-89-5H11	19.5	0.0	3.0	3.3	4.0	14.5
8668	RZM Z731H11	21.0	0.0	3.5	3.5	4.0	19.4
8939	RZM Y769H31	17.0	0.0	3.8	3.5	4.0	19.2
Mean		18.6	1.1		3.1	3.5	27.1
LSD (.05)		3.6	3.6		8.0	6.0	19.1
C.V. (%)		14.0	242.8	18.4	19.2	18.1	50.5
F value		1.8**	6.3**		7.1**	7.4**	6.3**

See notes for B1299.

October. Living plants counted 08 July 1999. The highest level of survival and best appearance in late resistance from WB97 and WB242 are highly rhizomania susceptible, when crossed to Rz, give a higher than (October 1998) are r = -.52**, -.74**, -.88**, and .03, respectively. Stand counts made post thinning in season again appeared to be associated with resistance to rhizomania from Beta maritima thru R22 (C50 & expected level of resistance to rhizomania (see P811 and P814 in this test and full-sib lines P807B and Coefficients of correlation for % Living vs. Appearance scores for 5/13, 6/11 & 7/8 and Stand Counts C51), e.g., lines R522, C67, & 8927-30. Although CP01 and CP02 that segregated for powdery mildew P808B in test B1299).

EVALUATION OF PAIR CROSSES (FULL SIB) FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99 TEST B1299.

Planted: September 24, 1998 Not harvested for yield 128 entries x 2 reps., sequential 1-row plots, 13 1/2 ft. long

	Description	Stand Count	% Bolting	Appe	Appearance	Score	Living Plants
		No.	06/10	05/12	06/11	01/08	જી∣
Spre	Spreckels, 9-98, L1162401	9					٦.
Spr	9-98, X78	7					7
Bet	776R.70	2		•			Ö
1997	7	9	•	•			Ή.
RZM	RZM-%S R322R4,(C51)	17.5	25.8	2.0	1.5	1.0	8.89
RZM	RZM 7926	5	٠	٠	•		ъ.
Inc	Inc. U86-37	٦.	0.0	•	•		
RZM	RZM Y775	4.	•	•	•	•	•
146PX =	$RZM R746PX = C37*3 \times R22 (ab 5)$						
RZM		5	•	•	•	•	52.7
		18.0	0.0	2.5		•	•
			•	3.0	3.5	3.5	40.9
		18.5	0.0	3.0	•	•	13.6
		•	•	•	•	•	47.0
		17.0	0.0	1.5	2.0	2.0	ı.
		•	•	•	•	•	19.6
		4.	•	•	•	•	1.
RZM R753PX =	$C37*4 \times R22 \text{ (qh 5)}$						
RZM	R753PX	9	•	•	•		8.3
		9	•	•	•	•	1.
		22.5	0.0	2.5	2.5	3.0	38.0
		Η.	•	•	•	•	4.
		7.	•	•	•	٠	9
		4.	•	•	•	•	•
		6.	•	•	•	•	15.9
		4.	•	٠	•	•	4.

TEST B1299. EVALUATION OF PAIR CROSSES (FULL SIB) FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99

No. $06/10$ $05/12$ $06/11$ $07/08$ = $RZM\ R753PX = C37*4\ x\ R22\ (gh\ 5)$ (cont.) 17.5 0.0 2.5 2.5 3.5 16.0 0.0 4.0 4.0 4.0 4.0 16.0 0.0 2.5 3.0 3.0 3.5 14.0 0.0 2.5 3.5 3.5 14.0 0.0 2.5 3.5 3.5 14.0 0.0 2.5 3.5 3.5 14.0 0.0 2.5 3.5 3.5 14.0 0.0 2.5 3.5 3.5 18.0 0.0 3.0 3.0 3.5 18.1 0.0 0.0 3.0 3.0 3.5 18.5 0.0 3.0 3.0 3.5 18.5 0.0 3.0 3.0 3.5 18.5 0.0 3.0 3.0 3.5 18.5 0.0 3.0 3.0 3.5 18.5 0.0 3.0 3.0 3.5 18.5 0.0 3.0 3.0 3.5 18.5 0.0 3.0 3.0 3.5 18.5 0.0 3.0 3.0 3.0 18.5 0.0 0.0 3.5 3.5 18.6 0.0 0.0 3.5 3.5 18.7 0.0 0.0 3.5 3.0 3.0 18.8 0.0 0.0 3.5 3.0 3.0 18.9 0.0 0.0 3.5 3.0 3.0 18.0 0.0 0.0 3.0 3.0 4.0 18.0 0.0 0.0 3.0 3.0 4.0 18.0 0.0 0.0 3.0 3.0 4.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 4.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 3.0 3.0 3.0 18.0 0.0 0.0 0.0 3.0 3.0	Variety	Description	Stand	% Bolting	Appe		Score	Living Plants
			No.	ဖြ	2/	06/11	 	oko
RZM K753PX 1175 0.00 2.5 2.5 3.0 3.0 3.0 3.0 3.0 3.0 3.0 3.0 3.0 3.0	11	C37*4 x R22 (gh 5)	1					
0 16.0 0.0 4.0 4.0 4.0 15. 2 20.0 0.0 4.0 4.0 4.0 15. 2 20.0 0.0 2.5 3.6 4.0 12. 1 4.0 0.0 2.5 3.6 4.0 12. 1 19.0 0.0 4.0 4.5 4.5 4.5 6. 1 18.0 0.0 3.0 3.0 3.5 4.1 1 19.0 0.0 4.0 3.6 4.5 6.0 1 19.0 0.0 3.0 3.0 3.5 3.5 3.1 1 10.0 0.0 3.0 3.0 3.5 3.5 3.1 1 10.0 0.0 3.0 3.0 3.0 3.5 3.5 3.1 1 10.0 0.0 3.0 3.0 3.5 3.5 3.1 1 10.0 0.0 3.0 3.0 3.5 3.5 3.1 1 10.0 0.0 3.0 3.0 4.0 4.5 14. 1 10.0 0.0 3.0 3.0 4.0 4.5 14. 1 10.0 0.0 1.5 1.5 2.5 3.0 3.0 4.0 1 10.0 0.0 1.5 1.5 2.5 3.0 3.0 1 10.0 0.0 1.5 1.5 2.0 5.0 1 10.0 0.0 3.0 3.0 4.0 5.0 1 10.0 0.0 1.5 1.5 2.0 5.0 1 10.0 0.0 3.0 3.0 4.0 5.0 1 10.0 1.5 1.5 2.0 5.0 1 10.0 1.5 1.5 2.0 5.0 1 10.0 1.5 1.5 2.0 5.0 1 10.0 1.5 1.5 2.5 3.5 3.0 1 10.0 0.0 1.5 1.5 2.0 5.0 1 10.0 1.5 1.5 2.5 3.5 3.0 1 10.0 0.0 1.5 1.5 2.5 3.5 3.0 1 10.0 0.0 1.5 1.5 2.0 5.0 1 10.0 1.5 1.5 2.0 5.0 1 10.0 1.5 1.5 2.5 3.5 3.5 1 10.0 0.0 1.5 1.5 2.5 3.5 1 10.0 0.0 1.5 1.5 2.5 3.5 1 10.0 0.0 1.5 1.5 2.5 3.5 1 10.0 0.0 1.5 1.5 2.5 3.5 1 10.0 0.0 1.5 1.5 2.5 3.5 1 10.0 0.0 1.5 1.5 2.5 3.5 1 10.0 0.0 1.5 1.5 2.5 3.5 1 10.0 0.0 1.5 1.5 2.5 1 10.0 0.0 1.5 2.5 1 10.0 0.0 1.5 2.5 1 10.0 0.0 1.5 2.5 1 1	853 - 9	RZM R753PX	۲.	•	•	•	•	· ·
1 20.0 0.0 3.0 3.0 3.5 3.0 3.5 3.0 3.0 3.5 3.0 3.0 3.0 3.0 3.0 3.0 3.0 3.0 3.0 3.0	-10		9	•	•	•	•	ر س
20.5 0.0 2.5 3.0 4.0 12. 14.0 0.0 2.5 3.5 4.5 14. 19.0 0.0 2.0 2.5 3.5 4.5 44. 19.0 0.0 4.5 4.5 4.5 6. 18.0 0.0 4.0 3.5 4.5 6. 18.0 0.0 3.0 4.0 26. 19.0 0.0 2.0 2.5 3.5 4.5 6. 19.0 0.0 3.0 4.0 26. 20.5 0.0 3.0 3.0 4.0 26. 20.5 0.0 3.0 3.0 4.0 26. 20.5 0.0 3.0 3.0 4.0 28. 20.5 0.0 3.0 3.0 4.0 28. 20.5 0.0 3.0 3.0 4.0 28. 20.5 0.0 3.0 3.0 4.0 28. 20.5 0.0 3.0 3.0 4.0 28. 20.5 0.0 3.0 4.0 4.5 14. 20.6 0.0 1.5 1.5 2.5 3.5 34. 20.7 0.0 1.5 1.5 2.0 3.0 4.0 28. 20.8 0.0 1.5 1.5 2.0 5.0 5.0 5.0 5.0 5.0 5.0 5.0 5.0 5.0 5	-11		0	٠	•	•	•	0
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4 19.0 0.0 2.0 2.5 4.5 28 18.0 0.0 4.5 4.5 4.5 6. 18.0 0.0 4.5 4.5 4.5 6. 18.0 0.0 4.5 4.5 4.5 6. 18.0 0.0 4.0 3.5 4.0 28 18.0 0.0 3.0 3.0 4.0 28 20.5 0.0 3.0 3.5 3.5 3.1 20.5 0.0 3.0 3.5 3.5 3.1 20.5 0.0 3.0 3.5 3.5 3.1 20.5 0.0 3.0 4.0 4.5 5.0 29 21.0 0.0 3.0 4.0 4.5 14. 22.0 0.0 1.5 2.5 3.5 3.5 3.6 3.0 4.3 23.0 0.0 0.0 1.5 1.5 2.5 3.5 3.6 3.0 4.3 24.0 0.0 0.0 1.5 1.5 2.0 5.0 5.0 5.0 5.0 5.0 5.0 5.0 5.0 5.0 5	-13		4.	•		•	•	÷.
5 5 6 6 7 18.0 0.0 4.5 4.5 4.5 6. 18.0 0.0 18.0 18.0 26. 19.0 26. 19.0 19.0 19.0 19.0 19.0 19.0 19.0 19.	-14		σ.	•	•	•	•	æ
6 6 6 6 6 6 7 13.0 0.0 13.0 3.0 2.5 14.0 26. 15.0 16.0 0.0 3.0 3.0 3.0 2.8 13.5 0.0 3.0 3.0 3.5 3.5 3.5 3.0 3.5 3.5 3.0 3.5 3.5 3.0 3.5 3.5 3.0 3.5 3.5 3.5 3.0 3.5 3.5 3.0 3.5 3.5 3.5 3.0 3.5 3.5 3.5 3.0 3.5 3.5 3.5 3.0 3.5 3.5 3.5 3.0 3.5 3.5 3.5 3.0 3.5 3.5 3.5 3.0 3.5 3.5 3.5 3.0 3.5 3.5 3.5 3.0 3.5 3.5 3.5 3.0 3.5 3.5 3.0 3.5 3.5 3.5 3.0 3.0 3.0 3.0 3.0 3.0 3.0 3.0 3.0 3.0	-15		ω.	•	•	•	•	•
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	-16		e,	•	•	•	•	9
$13.5 \qquad 0.0 \qquad 3.0 \qquad 2.5 \qquad 3.5 \qquad 3.5$ $20.5 \qquad 0.0 \qquad 3.0 \qquad 3.5 \qquad 3.5 \qquad 3.1$ $18.5 \qquad 0.0 \qquad 3.0 \qquad 3.5 \qquad 3.5 \qquad 3.1$ $12.0 \qquad 0.0 \qquad 2.5 \qquad 2.5 \qquad 3.5 \qquad 3.1$ $12.5 \qquad 0.0 \qquad 2.5 \qquad 2.5 \qquad 3.5 \qquad 45$ $12.5 \qquad 0.0 \qquad 4.0 \qquad 4.0 \qquad 4.5 \qquad 14$ $18.0 \qquad 0.0 \qquad 1.5 \qquad 2.5 \qquad 3.0 \qquad 3.0$ $18.5 \qquad 0.0 \qquad 4.0 \qquad 4.0 \qquad 4.0 \qquad 4.0$ $16.0 \qquad 0.0 \qquad 1.5 \qquad 1.5 \qquad 2.0$ $20.0 \qquad 0.0 \qquad 1.5 \qquad 1.5 \qquad 2.0$ $20.0 \qquad 0.0 \qquad 1.5 \qquad 1.5 \qquad 2.0$ $20.0 \qquad 0.0 \qquad 3.0 \qquad 3.0 \qquad 4.0 \qquad 2.5$ $16.0 \qquad 0.0 \qquad 3.0 \qquad 3.0 \qquad 4.0 \qquad 2.0$ $17.5 \qquad 0.0 \qquad 3.0 \qquad 3.0 \qquad 4.0 \qquad 3.0$ $20.5 \qquad 0.0 \qquad 4.0 \qquad 4.0 \qquad 4.0 \qquad 2.0$ $20.5 \qquad 0.0 \qquad 1.0 \qquad 1.0 \qquad 1.5 \qquad 3.0$ $20.5 \qquad 0.0 \qquad 1.0 \qquad 1.0 \qquad 1.5 \qquad 5.0$ $20.5 \qquad 0.0 \qquad 1.0 \qquad 1.0 \qquad 1.5 \qquad 5.0$ $20.5 \qquad 0.0 \qquad 1.0 \qquad 1.0 \qquad 1.5 \qquad 5.0$ $20.5 \qquad 0.0 \qquad 1.0 \qquad 1.0 \qquad 1.5 \qquad 5.0$	-17		9	•	•	•	•	80
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$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	-21		8	•	•	•	•	Ŋ.
	-22		2	•	•	•		4.
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EVALUATION OF PAIR CROSSES (FULL SIB) FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, TEST B1299.

1998-99	
VALLEY,	
IMPERIAL	

RZM Y773PX = $F_2(C37 \times Y71(C))$ (gh 5) (cont.) RZM Y773PX 16.0 13.5 20.5 17.5 17.5 17.5 17.5 17.5 17.5 18.5
1(C)) (gh 5) (cont.) 16. 13. 20. 17. 17. 18.
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TEST B1299. EVALUATION OF PAIR CROSSES (FULL SIB) FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99

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Living	Plants		25.0	43.3		•	68.5	•	17.0	45.7	30.8	60.5		34.0	63.9	37.5	64.7	16.7	71.9	41.7	49.1	74.8	74.6		5.3	31.8
	07/08		. 4 . 0	3.5	1.5	2.0	1.5	3.0	•	2.5	•	•		•	1.5	•	•	•	•	•	•	•	•			2.5
	Appearance S 12 06/11			2.0	•	•	•	•	•	1.5	•	•		•	1.5	•	•	•	•		•	•	•		•	1.5
•	Appe 05/12		2.5	2.0	•	•	•	•		2.0				•	1.0	•	•	•	•	•	•	•	•		•	2.0
% . !	06/10	c	0.0	0.0	0.0	2.5	0.0	15.8	0.0	0.0	0.0	0.0		0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	•		0.0	2.1
Stand	No.			17.5	6	•	16.5	7.	•	4.	13.0	7.		14.5	15.5	۲,	ω.	9	16.0	8	18.5	•	5.		9	22.5
:	Description	$RZM Y771PX = 0.P. \times R22 \text{ (gh 5)}$	KGM I / IFA										$RZM Y772 = R80, R76 \times (C37 \times R22)$ (gh 5)												1997	RZM 7926aa x A
:	Variety		18/T = T	ı m I	- 4	ı S	9 -	L - 7	80 I	6 I	-10	-11	II	Y872 - 1	- 2	m I	- 4	- 5	9 -	- 7	& 1	ი 1	-10	Checks	US H11	8927

TEST B1299. EVALUATION OF PAIR CROSSES (FULL SIB) FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99

(cont.)
Count No.
(C913-70aa x R636)
P603) (~CP01)) (gh

EVALUATION OF PAIR CROSSES (FULL SIB) FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99 TEST B1299.

		(conc.)					
		Stand	℀				Living
Variety	Description	Count	Bolting	Appe	Appearance Score	Score	Plants
		No.	06/10	05/12	06/11	01/08	o-⊱
P808B-# = R778%	P808B-# = R778% x RZM P708B ((Y71 x P604) (\sim CP02)) (gh 10)	(gh 10)					
P808B- 2	RZM P708B x R778%	11.5	0.0	1.5	1.5	2.5	63.1
m I		18.5	0.0	4.0	4.0	4.0	5.4
4 -		18.0	0.0	3.5	3.0	3.5	30.5
- 7		18.5	0.0	1.0	1.5	1.5	56.9
Checks							
X867	RZM Y767 (C67)	18.5	0.0	1.5	1.0	1.0	73.1
X872	RZM-%S Y672 (C72)	18.0	0.0	1.5	1.0	1.5	67.6
US H11	1997	13.5	0.0	4.5	5.0	5.0	0.0
R522 (Sp)	RZM-8S R322R	17.0	14.7	1.5	1.0	1.0	82.4
Mean		16.7	0.7	2.7	2.6	3.2	37.1
LSD (.05)		4.9	5.1	1.3	1.2	1.4	32.8
C.V. (%)		14.8	345.4	24.3	23.9	22.8	44.7
F value		3.2**	3.6**	4.3**	e.0**	4.7**	3.5**

OTES:

Appearance scored on a scale of 1 to 5 where: 1 = very good canopy; 2 = good canopy and appearance often segregating; 3 = intermediate and variable; <math>4 = fair; and 5 = poor to mostly dead plants.

However, other factors such as plant vigor, cyst nematode infection, root rots, etc. could have influenced vigor, number of dead leaves, and dead plants. The assumption was that plant health and appearance was Appearance scored relative to the overall test at time and based upon canopy size, uniformity, color, mostly being influenced by reaction to rhizomania and rhizomania under high temperature conditions. appearance. Coefficients of correlation for % Living plants vs. Appearance scores for 5/12,6/11, & 7/8 and Stand Counts (October 1998) are r = -.60**, -.72**, -.87**, and 0.01, respectively. Stand counts made post thinning in Living plants counted 08 July 1999. October.

TEST B1399. EVALUATION OF MONOGERM S1 PROGENY FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, INPERIAL VALLEY, 1998-99

138 entries x 1-row plots,	: 1 or 2 replications, sequential 13 1/2 ft. long			Planted: Not harv	Planted: September Not harvested for yi	oer 24, 1998 : yield
Variety	Description	Stand	Appe	Appearance	Score	Living Plants
		No.	05/12	06/10	07/08	₩
Checks						
8835	7835mmaa x A	•	3.0	•	•	22.9
8838	X X	•	•	•	•	5.8
8848M	RZM 7848 (=C790 x 848)	21.0	•	•	3.0	26.1
8810M	RZM 7810NB	•	•	•	•	32.5
8818-1B	Inc. 6818B-1	19.5	3.0	2.5	4.0	10.4
8818-2B	Inc. 6818B-2		4.0	•	4.5	2.9
of	$6818 - \# S_1's = C790 \times R22 = C790 - 8$					
	Inc. 6818- 1mm (A, aa)	9.5	•	4.0	•	7.1
8818- 2(C)	6818- 2mm	10.5	3.5	4.5	5.0	0.0
8818- 6(C)	Inc. 6818- 6mm (A,aa)	17.5	•	•	•	8.8
8818-11 (C)	Inc. 6818-11mm (A, aa)	9.5	•	•	4.5	3.6
8818-12 (C)	Inc. 6818-12mm (A, aa)	19.0	•	3.5	4.5	18.4
8818-21 (C)	Inc. 6818-21mm (A,aa)	11.0	•	•	4.0	11.5
S_1 's of popn-818	$318 = C790 \times R22 = C790-8$					
- 1	RZM-%S 6818mm⊗	18.0	3.5	•	3.5	28.1
- 2		14.5	•	•		•
n ع		17.5	1.0	1.0	2.0	63.3
- 4		•	•	•		
ı S		•	•	•		•
9 -			•	•		41.0
- 7		5	•	•	•	•
80 I		•	•	•		•
၈ ၊		ъ	•	•	•	•
-10		8	•	•	•	•
-11		o.	•	•	•	•
-12		8.0	•	•		0.0

TEST B1399. EVALUATION OF MONOGERM S₁ PROGENY FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99

Varietv	Description	Stand	Acce	Appearance	S. O.	Living Plants
		No.	05/12		80/20	o⁄e
S_1 's of popn-808	$= C790 \times 808(C) = C890-#'s$					
8808 - 1	RZM-%S 6808mm⊗	ω.	•	•	•	0.0
1 2		9	•	•	•	
د ا		•	•	•		7.0
4 -		5.	•	•	5.0	
ا 5		17.0	4.0	3.5	4.5	14.3
9 1		18.5		•	4.5	5.3
8808 - 7	RZM-%S 6808mm⊗	•		•	•	5.3
80 I		7.	•	•	•	5.
<u></u> რ		ω	•	•	•	8
-10		20.0	2.5	2.5	4.0	17.4
-11		ი	•	•	•	7.
-12		9	•	•	•	•
8808 -13	RZM-%S 6808mm⊗	14.5	•	•	4.0	
-14		•		•	•	20.2
-15		•	•	•	•	9
-16		13.5	2.5	3.0	4.5	4.2
-17		٠	•	•	•	•
-18		•		•	•	41.4
8808 -19	RZM-%S 6808mm⊗	55.0	4.0	4.0	4.5	7.1
-20		no plants	· ·	ļ. !	1.1	ļ. Ī
-21		16.0	•	•		0.0
-22		•	•	•		•
-23		15.5	3.0	2.5	4.0	ά.
-24			•	•	•	•

EVALUATION OF MONOGERM \mathbf{s}_1 PROGENY FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99 TEST B1399.

(cont.)

Living Plants	ok∙	15.3	41.2	46.1	15.3	5.3	15.0		•	•	•	•	•	0.0	•	0.0	•	•		0.0	20.0	0.0	0.0	0.0	0.0	0.0
Score	01/08			2.0	•	4.5	4.5		5.0	5.0	5.0	5.0	5.0	5.0	5.0	•	•	•	•	5.0	•	5.0	5.0	5.0	5.0	5.0
Appearance S	06/10	2.5	2.0	1.0	2.0	3.0	3.0		•	4.0	•	3.0	•	4.0	•		•	•	•	4.0	•	4.0	4.0	5.0	4.0	4.0
Appe	05/12			2.0	•	•	•		4.0		5.0			5.0		4.0	4.0	4.0	4.0	4.0	4.0	4.0	5.0	•	4.0	4.0
Stand Count	No.	15.5	18.0	17.5	19.5	18.5	17.5		14.0					19.0		16.0	7.	5	ω.	16.0	·	16.0	4	0.6	16.0	
Description		Inc. U86-37	(C79-1	RZM R736 (C70-8Rz)	Inc. 6818B-1	Inc. 6818B-2	RZM 7810NB	O Indexing (1 rep)	7808-2mm⊗							7808-3mm⊗						7808-4mm⊗				
Variety		Checks 97-C37	R879	R836	9818-1B	9818-2B	8810M	S2's from Type-O	8808 -2-1	-2-2	-2-3	-2-4	-2-5	-2-6	-2-7	8808 -3-1	-3-2	-3-3	-3-4	-3-5	-3-6	8808 -4-1	-4-2	-4-3	7-7-	-4-5

TEST B1399. EVALUATION OF MONOGERM S1 PROGENY FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99

808 + 9-1	Variety	Description	Stand Count	Appe	Appearance (Score	Living Plants
TBOB-Jamm⊗ 7808-Jamm⊗ 13.0 4.0 4.0 5.0 RZM 6808⊗ 13.0 4.0 4.0 5.0 7808-8mm⊗ 17.0 4.0 4.0 5.0 7808-8mm⊗ 17.0 4.0 4.0 5.0 7808-9 18 5.0 4.0 4.0 4.0 5.0 7808-12mm⊗ 15.0 5.0 4.0 4.0 5.0 7808-12mm⊗ 5.0 4.0 4.0 4.0 5.0 7808-12mm⊗ 5.0 5.0 4.0 5.0 7808-12mm⊗ 5.0 5.0 4.0 5.0 7808-12mm⊗ 5.0 4.0 5.0 5.0 7808-12mm⊗<			No.	05/12	06/10	80/10	o,e
7808-4mm⊗ 13.0 4.0 4.0 5.0 1.0 1.0 5.0 4.0 5.0 1.0 5.	S2's from Type-	Indexing (1 rep)					
RZM 6808⊗ 13.0 4.0 5.0 4.0 5.0 7808-8mm⊗ 12.0 3.0 3.0 5.0 15.0 15.0 3.0 5.0 15.0 15.0 3.0 5.0 15.0 15.0 3.0 5.0 15.0 17.0 5.0 4.0 5.0 17.0 5.0 4.0 5.0 17.0 5.0 4.0 5.0 17.0 5.0 4.0 5.0 17.0 17.0 4.0 5.0 5.0 17.0 5.0 4.0 5.0 17.0 5.0 4.0 5.0 17.0 5.0 4.0 5.0 17.0 5.0 4.0 5.0 17.0 5.0 4.0 5.0 17.0 5.0 17.0 5.0 4.0 5.0 17.0 5.0	8808 -4-6	7808-4mm⊗	•	•	•	5.0	0.0
RZM 6808⊗ 13.0 4.0 4.0 3.0 5.0 0 15.0 17.0 1	-4-7		11.0		4.0	5.0	0.0
21.0 4.0 3.0 5.0 0.0 15.0 15.0 3.0 5.0 0.0 15.0 17.0 4.0 3.0 5.0 0.0 0.0 17.0 4.0 5.0 5.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0	8808 -7-1	RZM 68088	•		•	5.0	0.0
15.0 3.0 5.0 0 7808-8mm⊗ 17.0 4.0 5.0 5.0 0 7808-8mm⊗ 4.0 4.0 5.0 0 7808-9 18 5.0 0.0 0 7808-12mm⊗ 19.0 5.0 4.0 5.0 0 7808-12mm⊗ 19.0 5.0 4.0 5.0 0 7808-12mm⊗ 19.0 5.0 4.0 5.0 0 11.0 4.0 5.0 0.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 4.0 5.0 0 11.0 5.0 6.0 0 11.0 5.0 6.0 0 11.0 6.0 6.0 6.0 0 11.0 6.0 6.0 6.0 0 11.0 6.0 6.0 6.0 0 11.0 6.0 6.0 6.0 0 11.0 6.0 6.0 6.0 0 11.0 6.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6.0 6 11.0 6.0 6 11.0 6.0 6 11.0 6.0 6 11.0 6.0 6 11.0 6.0 6 11.0 6.0 6 11.0 6.0	-7-2		21.0	•	•	5.0	0.0
7808-8mm⊗ 12.0 12.0 12.0 12.0 17.0 12.0 17.0 17.0 17.0 17.0 1808-8mm⊗ 7808-8mm⊗ 7808-9 18 12.0 12.0 12.0 12.0 12.0 12.0 12.0 12.0	-7-4		15.0	•	•	5.0	0.0
12.0 5.0 4.0 5.0 29 17.0 2.0 2.0 3.0 29 17.0 4.0 4.0 5.0 3.0 29 21.0 4.0 4.0 5.0 0 21.0 4.0 4.0 5.0 0 21.0 4.0 5.0 0 21.0 5.0 0.0 5.0 0 20.0 5.0 4.0 5.0 0 20.0 5.0 4.0 5.0 0 20.0 19.0 3.0 4.0 5.0 0 20.0 14.0 5.0 5.0 0 20.0 14.0 5.0 5.0 0 20.0 15.0 5.0 4.0 5.0 0 20.0 15.0 5.0 4.0 5.0 0 20.0 5.0 4.0 5.0 0 20.0 5.0 4.0 5.0 0 20.0 5.0 4.0 5.0 0 20.0 5.0 4.0 5.0 0 20.0 5.0 4.0 5.0 0 20.0 5.0 4.0 5.0 0 20.0 5.0 4.0 5.0 0	8808 -8-1	7808-8mm⊗	7.		3.0	5.0	•
7808-8mm⊗ 4.0 2.0 2.0 3.0 2.9 7808-9mm⊗ 4.0 4.0 4.0 5.0 0 12.0 4.0 4.0 5.0 0 12.0 4.0 5.0 0 12.0 4.0 5.0 0 15.0 5.0 4.0 5.0 0 15.0 5.0 4.0 5.0 0 18.0 4.0 5.0 0 19.0 3.0 4.0 5.0 0 114.0 5.0 0 15.0 4.0 5.0 0 115.0 5.0 4.0 5.0 0 115.0 5.0 4.0 5.0 0 115.0 5.0 4.0 5.0 0 115.0 5.0 4.0 5.0 0 115.0 5.0 4.0 5.0 0 115.0 5.0 4.0 5.0 0 115.0 5.0 4.0 5.0 0 115.0 5.0 4.0 5.0 0 115.0 5.0 4.0 5.0 0 115.0 5.0 4.0 5.0 0 115.0 5.0 4.0 5.0 0	-8-2		12.0	•	4.0	5.0	•
7808-8mm% 4.0 4.0 4.0 5.0 21.0 4.0 4.0 5.0 21.0 4.0 5.0 6.0 21.0 4.0 5.0 6.0 6.0 6.0 6.0 6.0 6.0 6.0 6.0 6.0 6	US H11		17.0	•	2.0	3.0	6
21.0 4.0 4.0 5.0 0 12.0 4.0 4.0 5.0 0 12.0 4.0 3.0 5.0 0 15.0 5.0 0.0 0 15.0 5.0 4.0 5.0 0 20.0 5.0 4.0 5.0 0 19.0 4.0 5.0 0 14.0 5.0 4.0 5.0 0 15.0 5.0 4.0 5.0 0 15.0 5.0 4.0 5.0 0 15.0 5.0 4.0 5.0 0 19.0 5.0 4.0 5.0 0 19.0 5.0 4.0 5.0 0 19.0 5.0 4.0 5.0 0 19.0 5.0 4.0 5.0 0 19.0 5.0 4.0 5.0 0 19.0 4.0 5.0 0	8808 -8-4	7808-8mm⊗	•	•	•	5.0	•
12.0 4.0 3.0 5.0 0.0 7808-9 18 5.0 3.0 5.0 0.0 15.0 5.0 4.0 5.0 0.0 20.0 5.0 4.0 5.0 0.0 18.0 4.0 5.0 0.0 19.0 3.0 4.0 5.0 0.0 20.0 4.0 5.0 0.0 14.0 5.0 4.0 5.0 15.0 5.0 4.0 5.0 15.0 5.0 4.0 5.0 19.0 5.0 4.0 5.0 19.0 5.0 4.0 5.0 19.0 5.0 4.0 5.0 19.0 4.0 5.0 19.0 4.0 5.0	-8-5		•	•	•	5.0	•
7808-9 18 5.0 5.0 5.0 6.0 15.0 15.0 15.0 15.0 15.0 15.0 15.0 16.0 18.0 19.0 19.0 19.0 19.0 19.0 19.0 19.0 19	-8-7		ς.	•	•	5.0	•
5.0 4.0 5.0 0 15.0 20.0 5.0 4.0 5.0 0 18.0 4.0 5.0 0 18.0 4.0 5.0 0 19.0 3.0 4.0 5.0 0 20.0 4.0 5.0 0 14.0 5.0 4.0 5.0 0 15.0 4.0 5.0 0 15.0 5.0 4.0 5.0 0 19.0 5.0 4.0 5.0 0 19.0 4.0 5.0 0 19.0 4.0 5.0 0 19.0 4.0 5.0 0	8808 -9-1	7808-9 18			•	0.0	
15.0 5.0 4.0 5.0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	-9-2		5.0	•	•	5.0	0.0
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	-6-3		S		•	5.0	0.0
18.0 4.0 4.0 5.0 0 19.0 3.0 4.0 5.0 0 20.0 4.0 5.0 0 14.0 5.0 4.0 5.0 0 15.0 4.0 5.0 0 15.0 4.0 5.0 0 20.0 5.0 4.0 5.0 0 19.0 5.0 4.0 5.0 0 19.0 4.0 5.0 0 19.0 4.0 5.0 0	-9-4		0	•	•	5.0	0.0
$19.0 \qquad 3.0 \qquad 4.0 \qquad 5.0 \qquad 0$ $20.0 \qquad 4.0 \qquad 4.0 \qquad 5.0 \qquad 0$ $14.0 \qquad 5.0 \qquad 4.0 \qquad 5.0 \qquad 0$ $15.0 \qquad 5.0 \qquad 4.0 \qquad 5.0 \qquad 0$ $20.0 \qquad 5.0 \qquad 4.0 \qquad 5.0 \qquad 0$ $19.0 \qquad 5.0 \qquad 4.0 \qquad 5.0 \qquad 0$ $19.0 \qquad 4.0 \qquad 5.0 \qquad 0$ $19.0 \qquad 4.0 \qquad 5.0 \qquad 0$	-9-5		œ	•	•	5.0	0.0
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	9-6-		6	•	•	5.0	0.0
	7-6-		0	•	•	5.0	0.0
$15.0 5.0 4.0 5.0 0$ $7808-12mm \otimes $	-9-11		4	•	•	5.0	0.0
$-12-1$ $7808-12mm\otimes$ 9.0 5.0 4.0 5.0 0 $-12-3$ 20.0 5.0 4.0 5.0 0 $-12-4$ 19.0 4.0 4.0 5.0 0 $-12-5$	-9-12		Ŋ	•	•	5.0	0.0
20.0 5.0 4.0 5.0 0 19.0 4.0 4.0 5.0 0 18.0 4.0 4.0 5.0 0	8808 -12-1	7808-12mm⊗	6	•	•	5.0	0.0
19.0 4.0 4.0 5.0 0 18.0 4.0 5.0 0	-12-3		0	•	•	5.0	0.0
18.0 4.0 4.0 5.0 0	-12-4		0	•	•	5.0	0.0
	-12-5		æ	•	•	5.0	0.0

0.0

5.0

4.0

5.0

13.0

-12-6 -12-7

EVALUATION OF MONOGERM \mathbf{s}_1 PROGENY FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99 TEST B1399.

Varietv	Description	Stand Count	Appe	Appearance 8	Score	Living Plants
		No.	05/12	1	01/08	& I
S ₂ 's from Type-O Indexing (1	indexing (1 rep) (cont.)					
	7808-13mm⊗	14.0		•	4.0	21.4
-13-2		18.0	3.0	3.0	5.0	
-13-3		19.0	•	•	4.0	15.8
-13-4		•	•	•	4.0	
-13-5		•	4.0	•	5.0	
-13-6		3.0	4.0	3.0	3.0	66.7
8808 -16-1	7808-16mm⊗	10.0	4.0	4.0	5.0	
-16-2		12.0	4.0	4.0	5.0	0.0
-16-3		•	4.0	4.0	•	
-16-4		14.0	4.0	4.0	5.0	•
-16-5		3.0	2.0	4.0	5.0	0.0
-16-6		0.9	2.0	4.0	5.0	0.0
-16-7		12.0	•	4.0	5.0	0.0
Checks 8835	7835mmаа х А	22.0	•		2.0	31.8
8838	×	22.0	5.0	3.0	4.0	13.6
8848	RZM 7848	19.0	•	•		52.6
S_1 's from C890-7 ((SES)					
8818-5(C)	Inc. 6817mm (A, aa)	13.0	4.0	4.0	4.0	0.0
8817-1	RZM-8S 6817mm⊗	17.0	4.0	•	4.0	0.0
8817-2		19.0	•	3.0		21.1
8817-3		•	•	•	3.0	56.3
8817-4		•	3.0	•	•	•
8817-5		15.0	•	•	4.0	20.0

TEST B1399. EVALUATION OF MONOGERM S₁ PROGENY FOR RESISTANCE TO RHIZOMANIA UNDER HIGH TEMPERATURES, IMPERIAL VALLEY, 1998-99

(cont.)

Living Plants	11.1 20.0 28.6 7.7 	0.000	0.0 0.0 7.1 8.7	0.0 0.0 0.0
Score 07/08	4 4 4 4 1 0 0 0 0 0 1 0 0 0 1 0 0 0 1 0 0 0 0 0	0 0 0 0 0 0 0 0	0.0. 4.4 0.0 0.4	
Appearance 12 06/10	W 4 W 4 I W O O O O I O	4 4 4 0	6 4 4 6 0	4 4 6 4 0 . 0 0 . 0
Appe 05/12	w w 4 w 1 w 0 0 0 0 1 0	4 7 7 8 0 . 0 0 0 . 0	9.0 0.0 0.0	w v w 4
Stand Count No.	18.0 10.0 14.0 13.0 no plants 9.0	19.0 10.0 20.0 18.0	19.0 12.0 14.0 23.0	19.0 19.0 20.0 16.0
Description	(R04) RZM-%S 6815mm⊗	(WB151) RZM-%S 6819mm⊗	C890-10 (WB169) RZM-%S 6820mm⊗	<u>1 (WB258)</u> RZM-%S 6821mm⊗
Variety	S1's from C890-5 8815-1 -2 -3 -4 -5	S ₁ 's from C890-9 8819-1 - 2 - 3	S ₁ 's from C890-1 8820 - 1 - 2 - 3	S ₁ 's from C890-11 8821 - 1 - 2 - 3 - 4

See notes for tests B1199 and B1299.

EVALUATION OF HERBICIDE TRANSGENIC HYBRIDS FOR YIELD, IMPERIAL VALLEY, CA., 1998-99 TEST B1499.

Planted: October 23, 1998 Harvested: June 13, 1999 6 entries x 8 reps., RCB 4-row plots, 24 + 3 ft. long

		Acre Yield)ld		Beets/	Root	Clean	
Variety	Description	Sugar	Beets	Sucrose	100	Rot	Beets	NO3-N
		I.bs	Tons	%	No.	%	%	Mean
Checks Rifle	Spreckels, 9-98, L1162401	7843	25.50	15.35	158	0.0	94.2	125
B4776R	Betaseed 4776R.7653 (3-27-98)	7597	24.94	15.21	147	0.0	95.3	128
Roundup-ready								
HM 115RR	Hilleshog Round-up ready	8784	30.06	14.63	157	0.0	95.4	64
HM 117RR	Hilleshog Round-up ready	7055	25.01	14.11	122	0.2	93.1	49
HM 116RR	Hilleshog Round-up ready	6348	23.91	13.29	150	0.5	93.1	81
Liberty-link								
8CG9372LL	Betaseed Liberty-link	7609	23.76	16.02	168	0.0	94.5	95
Mean		7539.4	25.53	14.77	150.3	0.1	94.3	93.4
LSD (.05)		501.9	1.30	0.55	10.9	0.3	1.1	25.8
C.V. (%)		9.9	5.03	3.65	7.2	301.2	1.2	27.2
F value		21.6**	26.06**	25.97**	16.7**	2.6*	6.7**	**6.6

EVALUATION OF HERBICIDE TRANSGENIC HYBRIDS FOR YIELD, IMPERIAL VALLEY, CA., 1998-99 TEST B1499.

(cont.)

									c	ע	؈	4	0.5NS
Impur.	Value	12686	12881		12543	12012	12255	12604	0	12490.9	1316.	10.4	
NH ₂ -N	шdd	546	625		585	570	533	909	ה ה	0.770	97.8	16.7	1.1NS
Potassium	wdd	2599	2317		2432	2173	2303	2272	0.00	2349.5	234.4	8.6	3.3*
Sodium	wdd	286	330		258	334	409	333	9	0.470	87.2	26.5	2.9*
Known SugarLoss	<u>1bs/a</u>	996	959		1122	899	871	968	, ca	1.706	74.9	7.8	12.2**
Recover. Sugar	& 	87.6	87.2		87.1	87.2	86.2	88.2	0	5.70	1.5	1.7	1.5NS
Recover. Sugar	lbs/t	269	265		255	246	229	283	0 7 7 0	6.107	12.2	4.7	19.6**
Recover. Sugar	<u>lbs/a</u>	6877	6638		7662	6157	5476	6713	2 7 2 2	0.000	528.0	7.9	15.8**
Variety		Checks Rifle	B4776R	Roundup-ready	HM 115RR	HM 117RR	HM 116RR	Liberty-link 8CG9372LL	W G G	Lidaii	LSD (.05)	C.V. (%)	F value

Round-up ready entries sprayed 1-14-99 with 1 qt/a Round-up Ultra. Liberty-link entry sprayed 1-4-99 establishment due to flee beetles and strong winds. This trial was replanted in October, resulting in lower weeding was required in Round-up and Liberty treated plots. Original planting in September had poor stand Otherwise plot was hand weeded and no conventional herbicides were used. Little yields than the Area 5 Coded Mid-harvest trial. with 28 ou/a Liberty. NOTES:

180 entries x 3 replications, sequential 2-row plots, 12 ft. long

Not harvested for yield

		Stand	BSDF	LP	CRT
Variety	Description	Count ¹	2nd ¹	9/22/992	9/14/993
		No.	Score	Score	Score
HYBRIDS					
US H11	Resistant check	20	4.0	4.3	4.3
WS-PM9	HM-WS-PM9, 4-18-95	23	4.0	4.0	2.0
B4776R	1-19-99	23	5.0	5.7	7.0
B4035R	Betaseed,	22	5.0	5.7	6.7
B4419R	1-19-99	22	4.0	4.3	5.0
B4430R	L4430.8052, 3-10-99	25	4.3	5.0	6.7
SS-432R	Spreckels, 2-8-99	23	4.0	4.3	5.7
Rizor	Spreckels, 2-8-99	23	5.7	6.3	7.3
Diff.	Smarakala 2.8-00	25	5.7	6.0	7 7
Rifle	Spreckels, 2-8-99	24		6.0	7.7
SS-NB7R	Spreckels, 3-3-98		4.7	5.3	6.3
SS-778R	X782402, 3-3-98	26	4.0	3.3	4.3
Monohikari	Seedex, 2-18-97	24	5.0	5.7	7.3
CR812H50	C790-15CMS x RZM CR712	22	4.7	4.7	5.7
CR813H50	C790-15CMS x RZM CR713	24	4.0	4.7	5.0
R876-89-5NBH50	C790-15CMS x RZM-% C76-89-5	24	4.3	4.7	5.7
R876-89-5H50	C790-15CMS x RZM-% C76-89-5	22	5.0	4.7	6.0
R576-89-18H50	$C790-15CMS \times C76-89-18$	24	4.7	5.3	6.0
R776-89-5H8	F82-546H3 x C76-89-5	24	4.3	5.0	6.0
R778H8	F82-546H3 x C78	21	4.3	4.3	4.7
R878%H50	C790-15CMS x RZM C78	23	4.3	4.0	4.7
R878H50	C790-15CMS x C78	22	4.7	4.7	4.7
R882H50	C790-15CMS x C82	17	4.7	5.0	6.7
R882H27	C831-4HO x C82	20	5.0	5.3	7.3
R882H37	C306/2CMS x C82	20	4.3	4.7	6.3
US H11	Posistant short	20	4.0	5 0	
Y769H8	Resistant check F82-546H3 x C69	20	4.0	5.0	6.0
Y769H39	C762-17CMS x C69	20	4.3	4.7	6.0
		23	4.0	4.0	4.3
Y869H50	C790-15CMS x C69	24	4.0	4.3	5.0
Y869H15-1B	6818-1Baa x C69	22	4.3	4.3	6.0
Y869H5	C833-5aa x C69	23	4.3	5.0	6.0
Y869H15-2B	6818-2Baa x C69	25	4.3	5.0	5.7
Y869H27	C831-4CMS x C69	23	4.3	4.0	5.0
3060U4E	0967 1040 060	0.6			
Y869H45	C867-1CMS x C69	24	4.0	4.3	4.7
Y869H46	7869-6HO x C69	26	4.3	4.7	5.3
Y869H18	7818HO x C69	26	4.3	4.7	6.0
Y869H29	C829-3aa x C69	22	4.3	5.0	6.0

Variety	Description	Stand Count ¹	BSDF 2nd ¹	LP 9/22/99 ²	CRT 9/14/99 ³
		No.	Score	Score	Score
HYBRIDS (cont.)				
<u> </u>	7835aa x C69	23	4.3	5.0	5.7
Y869H38	7838aa x C69	23	4.0	4.3	5.3
Y869H69	7869aa x C69	25	4.3	4.3	5.0
Y869H37	C306/2CMS x C69	25	4.3	4.3	5.3
Monohikari	Susceptible check	24	5.3	5.7	7.0
US H11	Resistant check	23	4.0	3.7	4.3
R879H50	C790-15CMS x C79-1Rz	23	4.0	4.7	5.3
R836H50	C790-15CMS x C79-8R22	24	3.7	4.3	5.0
R854H50	C790-15CMS x RZM R754	25	4.0	5.0	6.0
Y867H50	C790-15CMS x RZM C67	25	4.7	4.3	6.3
Y872H50	C790-15CMS x RZM-% C72	27	4.3	4.3	5.3
Y873BH50	C790-15CMS x RZM Y773	25	3.7	4.0	5.0
Y875H50 Sp	C790-15CMS x Y775	23	4.0	4.7	5.3
Y875H50 Iso	C790-15CMS x RZM Y775	28	4.3	5.3	4.7
Z831H50	$C790-15CMS \times RZM Z25/Z30$	23	4.0	4.7	5.3
8924H50	C790-15CMS x 7924	25	4.0	4.7	5.7
8931H50	C790-15CMS x RZM 7931	26	4.7	5.0	5.3
8931H38	7838mmaa x RZM 7931	23	4.3	4.7	5.7
8932H50	C790-15CMS x 7932CT,	23	4.0	4.3	4.3
8932Н38	7838mmaa x 7932CT,	23	4.0	4.3	5.7
8932н69	6869mmaa x 7932CT,	23	4.0	4.0	4.7
HM-WS-PM9	HM-WS-PM9, 4-18-95	25	4.0	4.3	3.7
8935H50 Iso	$C790-15CMS \times R776-89-5H13$	26	4.0	4.7	5.7
8935H38	7838mmaa x R776-89-5H13	21	4.3	5.3	6.0
8936H50	C790-15CMS x R776-89-5H31	26	4.3	5.0	5.3
8937H50	$C790-15CMS \times R776-89-5H11$	25	4.3	4.7	5.0
8938H50	$C790-15CMS \times Z731H11$	24	4.0	4.7	5.7
8939H50	C790-15CMS x Y769H31	24	4.0	4.3	5.0
8926H50 Iso	C790-15CMS x RZM 7926	25			5.0
8926H50 Sp	C790-15CMS x RZM 7926	24			4.0
R709-1H50	C790-15CMS x CR R509A-1	23	4.0		5.0
R710H50	C790-15CMS x CR R509/10-#	21	4.3	4.7	6.0
MULTIGERM, O.	P. LINES				
US H11	Resistant check	26	4.0	4.3	4.7
97-US75	Inc. 268 (US75)	24	4.0	5.3	4.0
97-US22/3	Inc. Y009 (US22/3)	23	4.0	4.7	4.0
WS-PM9	HM-WS-PM9, 4-18-95	25	4.0	3.7	2.0

Variety	Description	Stand Count ¹	BSDF 2nd¹	LP 9/22/99 ²	CRT 9/14/99 ³
		No.	Score	Score	Score
MULTIGERM. O.I	P. LINES (cont.)				
97-SP22-0	Inc. SP7622-0	27	4.7	5.7	7.3
98-EL-02	RZM 94-RM-#s	24	4.7	5.3	6.7
98-EL-04	RZM 94-RM-#s	23	4.7	5.3	6.3
R576-89-18(Sp)	Inc. C76-89-18	22	4.3	5.0	6.0
R876-89-5NB	RZM-% C76-89-5NB	23	4.7	5.3	6.3
R878%	RZM C78	24	4.0	4.3	4.7
R878 (Sp)	Inc. C78	22	4.0	4.0	3.0
R880	RZM C80	22	4.7	5.3	6.0
R881	RZM R776,R781,	22	5.0	5.3	6.3
R882	Inc. C82	20	4.3	5.3	6.3
Y868	RZM Y768	23	4.7	5.0	5.7
Y869(Iso)	RZM C69	21	4.7	5.0	6.7
Y869 (Sp)	Inc. C69	24	4.7	5.0	5.7
P601	PMR P401	23	4.0	4.3	4.3
P811	RZM-PMR 6203-6208	24	4.0	4.7	5.7
P813	Inc. CP01	25	3.7	4.3	5.0
P814	Inc. CP02	25	4.0	4.3	5.3
R824	RZM C79-2/3, WB41/42	22	3.7	4.3	4.3
R835	RZM C79-7, SES	22	4.0	4.3	5.0
R836	RZM C79-8, R22	25	3.7	4.3	5.0
R879	RZM C79-1, Rz	22	3.7	4.3	3.7
US H11	Resistant check	20	3.7	4.7	4.0
R840	RZM R740 (C79#s)	24	4.3	5.0	5.0
R853	RZM-ER-% R653	25	4.3	4.7	5.3
R854	RZM R754	24	4.0	4.7	4.7
R726	RZM-ER R526, (C26)	26	4.0	5.0	5.3
R827	RZM R727A,B	21	4.7	5.0	6.7
Y866	RZM Y766	22	4.7	5.0	6.0
Y867	RZM C67	24	4.7	5.0	6.3
Y871	RZM Y771	23	4.3	5.3	6.0
Y872	RZM-% C72	22	4.0	5.0	6.3
Y872B	RZM C72	22	4.3	5.3	6.0
¥873	RZM-ER-% Y673	23	4.0	4.7	5.7
Y873B	RZM Y773	22	4.3	5.0	6.0
Y875(Iso)	RZM Y775	23	4.7	5.7	6.3
Y875 (Sp)	RZM Y775,,Y767	20	4.3	5.0	6.0

Variety	Description	Stand Count ¹	BSDF 2nd ¹	LP 9/22/99 ²	CRT 9/14/99 ³
		No.	Score	Score	Score
\	Aa POPULATIONS & LINES				
CR811	RZM CR09/CR10	19	5.0	5.7	6.7
CR812	RZM CR09/CR10	23	4.7	5.3	6.3
CR813	RZM CR713	23	5.0	5.3	6.0
WS-PM9	HM-WS-PM9, 4-18-95	22	4.0	4.3	3.0
WS 1119	III WO 2115 / 1 10 50				5.0
8932M	7932CTaa x A	24	4.0	4.3	4.3
Y869H30M	7 9 32 CT aa x C69	21	4.3	5.0	5.7
P812	RZM-PMR 6211-6217	23	4.0	4.3	4.0
Z831	RZM Z731-Z725aa x A	21	4.0	5.0	6.0
8924	RZM 7924aa x A	22	4.7	4.7	6.7
8931	RZM 7931aa x A	21	4.3	5.0	5.7
Y869H31	7931aa x C69	20	4.0	5.0	5.3
8935 (Iso)	RZM R776-89-5H13	23	4.7	5.7	6.7
0026	RZM R776-89-5H31	21	4.7	5.3	6.0
8936 8937	RZM R776-89-5H31	22	4.3	5.3	6.3
8937	RZM Z731H11	22	5.0	5.7	7.0
8939	RZM Y769H31	22	4.3	5.0	5.7
0939	R2H 1703H31		1.5	3.0	
8926(Iso)	RZM 7926	23	4.3	5.0	5.7
8926 (Sp)	$7931aa \times RZM 7926$	25	3.7	4.7	4.3
N724	Inc. N623,N624	19	4.0	4.7	5.3
77 47	Inc. 5747 (A,aa)	20	4.0	4.7	4.7
7005 6	T 8625 6	20	4.0	5.0	5.7
Z825-6	Inc. Z625-6 Inc. Z625-9	22	4.3	5.3	7.0
Z825-9 Z830-11	Inc. Z630-11	22	4.0	5.0	6.7
2830-11 8911-4-10M	RZM-ER-% 6911-4-10	22	4.0	4.3	3.3
8911-4-10M	RAM-ER 6 USIT 4 10				
8913-70	RZM-ER-% C913-70	22	4.3	5.0	6.7
8918-12	RZM-ER-% 6918-12	17	4.7	5.3	6.3
8925-19	Inc. 6925-19	20	4.3	5.3	6.7
8927-29	Inc. 6927-29	20	5.3	6.3	7.7
	Inc. 6927-30	20	4.7	4.7	6.0
8927-30	Inc. 6927-30	18	4.0	4.7	4.7
8927-33	Inc. 6927-33	17	4.3	6.0	6.3
8927-37 8929-41	Inc. 6929-41	18	5.3	6.3	7.3
8929-41	Inc. 6929 41		3.5	0.5	
8929-72	Inc. 6929-72	17	5.0	5.7	6.7
8929-102	Inc. 6929-102	23	5.0	5.0	6.7
8929-112	Inc. 6929-112	22	4.3	4.7	5.7
8929-114	Inc. 6929-114	20	4.7	4.7	6.3
0000 117	To - C000 115	18	4.3	5.0	6.3
8929-115	Inc. 6929-115	18	4.3	5.0	5.3
8929-133	Inc. 6929-133	10	7.5	3.0	2.5

Variety	Description	Stand Count ¹ No.	BSDF 2nd ¹ Score	LP 9/22/99 ² Score	CRT 9/14/99 ³ Score
				<u> </u>	<u>55525</u>
MULTIGERM,	Sf, Aa POPULATIONS & LINES (cont)			
8929-153	Inc. 6929-153	18	4.7	5.7	6.7
8929-154	Inc. 6929-154	21	4.7	4.7	6.7
8930-19	Inc. 6930-19	23	4.7	5.3	5.3
8930-39	Inc. 6930-39	20	4.3	4.7	5.0
8930-102	Inc. 6930-102	16	4.0	5.0	6.0
US H11	Resistant check	24	4.0	4.3	5.3
MONOGERM, S	f, Aa POPULATIONS & LINES				
7818%M	RZM-ER 5818 (C890-8,R22)	21	4.3	5.3	6.3
8818-1B	Inc. 6818-1B	21	4.0	5.0	6.3
8818-2B	Inc. 6818-2B	25	4.7	5.3	6.3
6546	Inc. F82-546 (C546)	18	4.0	5.0	5.7
6718	Inc. U83-718 (C718)	13	4.0	4.3	3.3
7864-14M	Inc. C864-14	20	4.0	5.0	5.7
8833-5H50	$C790-15CMS \times 5833-5$	26	4.0	4.7	5.7
8833H50	C790-15CMS x RZM,T-0 7833	28	4.0	4.3	5.0
6869 (Sp)	5869mmaa x A	25	4.0	4.3	5.3
8869	RZM 7869-#s	29	4.3	4.7	6.0
8890m	RZM 7890	23	4.3	4.7	5.7
8833	RZM,T-O 7833-#s	24	4.7	5.3	6.7
8836	T-O 7836-#s	24	4.7	5.3	7.0
8835	7835mmaa x A	24	4.0	5.0	6.0
8835H50	C790-15CMS x 7835	22	4.0	4.3	5.3
8838	7838mmaa x A	22	4.0	4.7	5.7
8838H50	C790-15CMS x 7838	24	4.0	4.7	5.7
8848M	RZM 7848	24	4.0	5.0	5.7
8810M	RZM 7810NB	24	4.0	4.7	4.7
8829-3	Inc. C829-3	24	4.7	5.3	5.7
8831-3	Inc. C831-3	24	4.7	5.3	6.3
8831-4	T-O C831-4-#s	22	5.0	5.0	6.3
8833-5	Inc. C833-5	24	4.7	5.0	6.0
8833-12	Inc. C833-12	23	4.7	5.0	6.7
7867-1M	Inc. T-0 C867-1 (CTR)	21	5.3	5.7	6.7
7869-6	T-0 6869-6 (barbed)	24	4.3	5.0	6.0
6762-17	Inc. 0762-17 (C762-17)	24	4.0	3.7	2.7
6562	Inc. F82-562 (C562)	18	4.0	4.0	5.0

 $^{^{\}rm 1}$ By Beet Sugar Development Foundation $^{\rm 2}$ By Dr. Lee Panella, USDA-ARS, Fort Collins $^{\rm 3}$ By Dr. Clyde Trupp

TEST 3199-2. EVALUATION OF PROGENY LINES FOR POWDERY MILDEW RESISTANCE, SALINAS, CA., 1999 (USDA entries)

21 entries x 4 reps, sequential 1-row plots, 11 ft. long

Planted: April 13, 1999 Not harvested for yield

		Stand				_	
Variety	Description	Count		Powdery			
		Mean	08/12	08/20	08/26	09/02	<u>Mean</u>
USDA entries							
US H11	Susc. check	18	4.8	7.0	7.5	7.3	6.6
Rizor	Spreckels,2-8-99	17	4.0	6.3	7.0	7.5	6.2
Rival	HH103, L1032406	16	4.5	6.8	7.3	8.0	6.6
B4430R	Betaseed 4430.8052, 3	-10-99					
		18	3.0	4.8	5.3	6.3	4.8
P811	RZM-PMR 6203-6208(C)	18	3.8	4.8	5.8	6.3	5.1
P812	RZM-PMR 6211-6217(C)	18	3.5	5.0	6.0	6.8	5.3
P813	Inc. 6201-6202(C)	17	3.8	5.0	5.8	5.8	5.1
P814	Inc. 6205-6206(C)	15	2.8	4.0	5.0	5.3	4.3
P601	PMR P401	17	3.0	4.5	5.0	4.8	4.3
P603	PMR P403	18	3.0	3.5	4.8	4.3	3.9
P604	PMR P404	17	3.0	3.8	4.8	4.8	4.1
8918-12	RZM-ER-% 6918-12	17	1.8	3.0	3.5	3.8	3.0
Y039	Inc. Y939 (C39)	15	2.8	4.0	4.8	4.3	3.9
Y869(Iso)	RZM Y769 (C69)	14	3.3	4.5	5.3	5.0	4.5
8939	RZM Y769H31	15	2.5	4.3	5.0	5.0	4.2
R878%	RZM R778%	17	3.0	5.0	6.3	6.3	5.1
B4776R	Betaseed, 1-19-99	18	3.8	5.8	7.0	6.8	5.8
Rifle	Spreckels, 2-8-99	16	4.3	6.5	7.8	7.3	6.4
SS-432R	Spreckels, 2-8-99	15	3.5	5.3	6.5	6.0	5.3
SS-778R	Spreckels, X782402, 9	-16-98					
ob //ok	opioeneze, consensa,	17	3.5	4.8	6.0	6.5	5.2
B4419R	Betaseed, 1-19-99	16	4.5	7.0	8.3	7.5	6.8
	•						
Mean		16.6	3.4	5.0	5.9	6.0	5.1
LSD (.05)		1.9	1.1	1.1	1.1	1.0	0.8
C.V. (%)		8.0	21.9	14.8	13.0	12.3	11.6
F value		3.3**	3.9**	9.6**	9.9**	11.3**	12.7**

Notes: P811, P812, P813, P814, P601, P603, P604 segregate for resistance to powdery mildew. Resistance was transferred to C37 from Beta maritima lines WB97 and WB242. On a plot basis, the PM ratings largely reflect the C37-type susceptible segregates.

ERWINIA/POWDERY MILDEW EVALUATION OF LINES AND POPULATIONS, SALINAS, CA., 1999 TEST 3499.

80 entries x 3 1-row plots, 1	80 entries x 3 reps., sequential 1-row plots, 17 1/2 ft. long						P. Ir.	Planted: A Inoc. Ecb: Scored Ecb:	April 13, July 14,	1999 1999 , 1999
Variety	Description	ď	Powderv	Mildew Score	Score	a	Stand	Harvest Count	Erwinia	Rating
		08/23	08/31	80/60	10/04	Mean	No.	No.		%H
Multigerm, open	open-pollinated									
US H11	113102 (resistant check)	7.0	7.0	7.7		7.4	28.3		7.9	81.5
E740	Inc. E840 (C40 susc. ck.)	8.3	8.7		8.7	8.7	31.0	30.3	79.3	14.3
97-US22/3	Inc. Y009 (US22/3)	•	6.7	7.3		7.1	31.3	31.7	6	ω.
97-US75	Inc. 268 (US75)	6.7	7.0	•	8.0	7.3	31.0	33.0	14.6	74.7
97-C37	Inc. U86-37 (C37)	7.0	7.3	7.3		7.4	5.	6	4.8	
R878% (Iso)	RZM R778%, (C78)		•			•	4.	4	•	Ή.
R878 (Sp)	Inc. R778, R778%	5.3	5.7	6.3	6.7	0.9	23.3	23.3	3.5	87.0
P601	PMR P401		•	•	•	•	4.	9	•	e.
R880	RZM R780, (C80)	4.7	•		7.3	5.8	•	24.3	1.7	4.
R882 (Sp)	Inc. R781, R776, R781-43,	4.7	5.0	6.3	6.7	5.7	25.0	•	3.8	90.9
R881 (Iso)	RZM R776, R781, R681,	•			7.0	0.9	•	22.7		ნ
	RZM-ER R576	5.3	•	•	7.3			•	8.9	4.
R781	RZM-ER R581	3.7		5.7	6.0	4.9		33.0	•	82.1
R770	RZM-ER R570	5.0	5.7	6.7	7.0	6.1	28.3	28.3	8.7	84.1
98-EL02	F_2 (C80 x smooth root)	5.3			•	6.7		Э.	6.2	90.1
98-EL04	F_2 (C80 x smooth root)		6.7	7.0	7.0	•	•	e.	7.2	
R879	(C79-1,	5.3				•	7.		•	2
R836	RZM R736, R743 (C79-8,R22)	7.7	8.0	8.0	8.7	8.1	29.7	31.0	18.2	75.3
R853	RZM-ER-%S R653	5.7				•	ი	6	•	4.
R854	RZM R754	6.3			•	•	2	ص	•	Ω.

ERWINIA/POWDERY MILDEW EVALUATION OF LINES AND POPULATIONS, SALINAS, CA., 1999 TEST 3499.

Varietv	Description	Δ	Powderv	Mildew	S. OTC	4	Stand	Harvest	ま い に の に	Rating
200		08/23	08/31	80/60	1-11	Mean	No.	No.	DI	8H
Multigerm, open-	Multigerm, open-pollinated (cont.)									
X873	RZM-ER-8S Y673	•	•			•	8	80	7.6	ω.
Y873B	RZM Y773	•	•			7.0	i.	ω.	3.1	ω.
US H11	113102	6.3	7.7	8.7	8.7	7.8	31.0	32.3	0.5	97.9
E740	Inc. E840	•	•	•		8.5	œ.	4.	54.1	0
R576-89-18 (Sp)	Inc. R476-89-18	4.0	•	•	•	•	œ.	8		7.
R876-89-5NB	RZM-%S R576-89-5NB	4.3	5.0	5.0	5.7	5.0	31.7	31.3	7.9	89.2
X866	RZM Y766	•	•	•	•	•	4.	w	•	5.
X868	RZM Y768	3.7	•	•	•	•	Η.	2	2.0	7.
(OSI) 69LX	RZM-ER Y569		•	•		•	Η.	e.	•	4
Y869 (Iso)	RZM Y769 (C69)	•	5.0	•	6.7	•	5	6	3.4	9
X869 (Sp)	Inc. Y769(C69)	5.0	5.7	0.9	•	5.9	29.0	31.3	•	97.8
X871		•	6.3	•	8.0	•	·	∺.	6.6	7.
X867	RZM Y767 (C67)	4.0		•		4.9	∞	4.	Η.	
X872	RZM-8S Y672	•		•	•	•	Η.	ω.	•	62.6
E740	Inc. E840	7.0	8.0	8.7	8.0	7.9	30.7	31.0	92.4	6.4
US H11	113102	6.0	•	•	•	•	2	2	•	84.5
Y872B	RZM Y772 (C72)	•	•	•		•	ω.	Η.	4	۵.
Y875 (Iso)	RZM Y775	•	•	0.9	6.3	•	0	2	•	
Y875 (Sp)	RZM Y775, Y773, Y772, Y767	5.3	5.7	0.9	•	5.9	31.0	32.7	8.0	80.2
R840	RZM R740 (C79-#s)	•	•	8.7	8.3	•	6	Η.	•	Ή.
R726 (C26)	RZM-ER R526	6.0	6.3	7.7	8.0	7.0	ο	33.0	•	7.
R827 (C27)	RZM R727A,B	•	•	•	•	•	30.7	e.	5.1	90.8

ERWINIA/POWDERY MILDEW EVALUATION OF LINES AND POPULATIONS, SALINAS, CA., 1999 TEST 3499.

(cont.)

Varietv	Description	Ä	Powdery	Mildev	Mildew Score	di	Stand	Harvest Count	Erwinia	Rating
		08/23	08/31	80/60	10/04	Mean	No.	No.	Id	H%
Multigerm, Sf,	S ^f , Aa populations									
8926 (Iso)	RZM 7926 (A,aa)	4.7	5.7	6.7	7.7		30.7	31.3	4.7	9
8926 (Sp)	7931aa × RZM 7926	4.7	0.9	6.7	6.7	0.9	30.7	31.7	4.7	87.3
8927	RZM 7926aa x A				6.7		29.7			88.0
7931	6931aa x 931(C)	4.3	5.7	0.9	6.7	5.7	31.0	31.7	2.0	90.5
8931	RZM 7931aa x A	•			7.0	•	31.0			90.1
8924	924				7.7		o.	•	6.3	85.7
2831	RZM Z731,Z730aa x A				•	•	6.		13.8	
CR811	RZM CR711 (CR09/10)		•	•	•	6.3	2		6.7	5.
CR812	RZM CR712	5.0	0.9	7.0	8.0	6.5	31.0	32.3	4.4	81.8
CR813	RZM CR713		•	•	•	•	7.		9 .5	o.
US H11	113102	6.7	7.0	7.7	8.7		œ.	31.7	9.8	2
E740	Inc. E840	8.0	8.7	0.6	8.7	9.8	30.0	30.0	82.9	12.2
P811	RZM-PMR 6203-#,6208-#(C)	•	4.7	5.3	•	•	ъ.	9		5.
P812	RZM-PMR 6211-#,6217-#(C)	•	•		•		ъ.	6		7 .
P813 (CP01)	Inc. 6201-#,6202-#(C)		•				ъ.	ъ.	•	ω.
P814 (CP02)	6205-#,6206-#	5.0	5.3	0.9	6.3	5.7	26.0	26.7	1.6	94.0
N724	Inc. N623,N624 (galls)	•	•		•		8	7.	•	4.
N730	Inc. N629,N630 (galls)						7.	9	ت	73.5
8932	7932CT,7201, aa x A	•	•			•	ο.	H.	H.	7.
8932Am	Inc. 7932CT,7201A	7.0	7.0	7.0	7.7	7.2	29.0	29.0	21.8	9.99
8932HO (M)	7204-7216CMS × A		•			•	ä	0	9	7.
8932H69	6869mmaa x A		•			•	o.	÷.	4.	6

ERWINIA/POWDERY MILDEW EVALUATION OF LINES AND POPULATIONS, SALINAS, CA., 1999 TEST 3499.

							Stand	Harvest		
Variety	Description	P	owdery	Powdery Mildew	Score	0	Count	Count	ja	Rating
		08/23	08/31	80/60	10/04	Mean	No.	No.	DI	%H
Multigerm, S ^f ,	Aa populations (cont.)									
8935 (Sp)	Inc. R776-89-5H13Aa	•	•	•	7.0	6.3		34.0		o.
	RZM R776-89-5H13	0.9	6.7	6.7	7.0	9.9	32.3	32.0	4.2	87.6
US H11	113102	7.3	•	•	•	7.9		2	2.8	•
E740	Inc. E840	8.7	0.6	0.6	8.0	8.7	29.7	29.7	85.4	7.8
8936	RZM R776-89-5H31	4.7	•		•	5.4	ο.	31.7	5.8	89.5
8937	RZM R776-89-5H11	3.7	•	•	•	5.2	31.3	31.0		7.
8638	RZM Z731H11	5.3	5.3	0.9	6.3	5.8	•	0	2.8	93.0
6863	RZM Y769H31	5.3	•	•	•	6.2	29.0	29.0		80.3
х 869н31	7931aa x Y769	4.3	•				უ.	8	•	•
7933	Inc. 6264-#(C)	•	5.7	6.3	7.3	6.2	31.7	H.	5.3	91.5
R710	CR-RZM R509-#, R510-#(C)	6.3	7.0	7.3		7.1	30.0	30.3	4.0	95.5
R709-1	CR-RZM R509A-1	•	0.9	7.3	8.0	6.8	œ.	÷.	4.2	91.4
R709-9	CR-RZM R509A-9	3.7	4.3	5.7	6.7	5.1	26.7	29.0	7.4	
2725	$Z625-\#(C)aa \times Z31(C)$	5.0	6.0	7.0	6.7	6.2	•	29.7		77.9
Z730	$Z630-\#(C)aa \times Z31(C)$	0.9	•				30.7	31.0	15.3	77.0
7747	Inc. 5747 (A,aa)	7.0	7.3	8.0	8.3	7.7	30.7	30.7	•	95.8
Mean		•	6.2		7.3	6.5	29.1	30.5	12.1	80.1
LSD (.05)		1.5	1.1	1.0	1.0	6.0	6.7	6.4	5.6	10.8
C.V. (%)		16.5		•	•	8.1	14.3	13.1	28.7	8.3
F value		4.6*	* 8.4**	8.1**	6.2**10	10.6**	1.3NS	1.4NS	86.9**	25.2**

ERWINIA/POWDERY MILDEW EVALUATION OF MULTIGERM, Sf, Aa PROGENY LINES, SALINAS, CA., 1999 TEST 3599.

40 entries x 1-row plots,	40 entries x 3 reps., sequential 1-row plots, 17 1/2 ft. long						Plan Inoc	Planted: April Inoculated Ecb: Scored Ecb: Nov	ti 13, 199 tb: July 1	.4, 1999 999
))
	4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	À	Dougle of the	7. Y	0,000		Stand	Harvest	000	0 0 0 0
Variety	Describution	08/23	08/31	09/08	-11	Mean	No.	No.	Score	% - VVI
Commercial	Hybrids									1
Rifle	Spreckels, 2-8-99	•	•	•	•	•		。	17.8	9
B4776R	Betaseed, 1-19-99	6.7	7.0	7.3	7.3	7.1	29.7	30.0	ω	73.5
US H11	Ecb. Resist. ck.	•	•	•	•	•		÷.	•	9
E740	Inc. E840, Ecb susc. ck.	•	•	•	•	•	œ ·	ij.	72.4	0
Rizor	Spreckels, 2-8-99	•	•				6	÷		
B4419R	Betaseed, 1-19-99	•	•	•	•	•	÷.	ς.	•	æ
SS-432R	Spreckels, 2-8-99	6.3	6.3	6.7	7.0	9.9	32.0	28.3	2.9	91.9
SS-778R	Spreckels, X782402, 9-16-98	•	•	•	•	•	。	1.	•	
B4430R	Betaseed 4430.8052, 3-10-99	•	•	•	•	•	4.	4.	2.	· 6
SS-NB7R	Spreckels, 3-3-98	•	•	•	•	•	ف	ö	4.	4.
B4035R	Betaseed, 7-10-97	6.3	7.0	7.7	7.0	7.0	29.7	31.7	18.6	72.7
Rival	HH103, L1032406, 3-18-97	•	•	•	•	•	· •	4	4.	4
es	of S_1 , MM , $S^{\mathcal{E}}$, Aa progeny lines									
8913-70	S 6913-	•	•		•	•	ö	$\vec{+}$	•	6
8918-12	RZM-ER-%S 6918-12	4.0	4.7	5.0	5.0	4.7	31.0	31.3	1.3	94.8
8918-21	RZM 7918-21	•	•		•	•	ά.	4.	•	თ
8911-4-10M	RZM-ER-%S 6911-4-10	•	•		•	•	œ.	7.	•	<u>o</u>
8925-19	Inc. 6915-19	•	•	•		•	о О	ω.	•	ъ.
Z825-6	Inc. Z625-6 (A, aa)	•	•	•	•	•	ж Э	ω.	•	ä
Z825-9	Inc. Z625-9 (A, aa)	4.7	5.0	5.7	0.9	5.3	28.3	30.3	23.8	65.8
Z830-11	Inc. Z630-11 (A,aa)	•	•	•	•	•	٠	ω.	•	4
E740	Inc. E840	•	•	•	•	•	œ.	7	•	
US H11	Ecb resist. ck.	7.3	7.3	7.3	8.0	7.5	33.0	32.3	4.2	85.8
8929-41		•	٠	•	•	•	5	ω ω	•	•
8929-72	Inc. 6929-72 (A,aa)	•	•	•	•	•	о О	.	14.9	5

(cont.)

Variety		Description	й	Powdery Mildew Score	Mildew	Score		Stand	Harvest Count	ERR-DI	ERR-8H
			08/23	08/31	80/60	10/04	Mean	No.	No.	Score	%
Increases	of S1/	MM, S ^f , Aa progeny lines	(cont.	?							
8929-102	Inc.	6929-102 (A,aa)		•	•	•	•	0	32.7	•	o,
8929-112	Inc.	6929-112 (A, aa)	5.7	5.7	7.0	6.7	6.3	29.3	31.0	0.7	6.96
8929-114	Inc.		4.7	•	6.3	5.7	•	ω.	31.3	20.8	6
8929-115	Inc.	6929-115 (A,aa)	4.0	•	•	6.0	•	ъ.	•	8.4	82.2
8929-133	Inc.	6929-133 (A,aa)		•	7.0	•	6.2	5.	27.0	9.2	79.2
8929-153	Inc.	6929-153 (A,aa)	5.3	0.9	•	6.3	6.1	29.3	•	•	5
8929-154	Inc.	6929-154 (A,aa)	•	4.7	5.7	•	4.9	4.	9	•	8.86
8930-19	Inc.	·	4.3	•	6.3	•	•	o.	30.0	6.3	7 .
8930-39	Inc.	6930-39 (A,aa)	4.7	•	•	•	•	œ.	о	•	
8930-102	Inc.	6930-102 (A, aa)	•	•	•	•	•	7.	7.	•	
8927-29	Inc.	6927-29 (A,aa)	4.7	5.3	0.9	5.3	5.3	26.7	28.0	1.3	7.76
8927-30	Inc.	6927-30 (A,aa)	•	•	•	•	•	œ.	ω.	•	•
8927-33	Inc.	6927-33 (A,aa)	•	•	•	•	6.4	ი		9.0	4
8927-37	Inc.			•	•	•	•	9		5.8	9
E740	Inc.	E840	0.6	0.6	0.6	8.0	8.8	30.0	31.7	73.3	21.1
US H11	Ecb 1	resist. ck.		•	•	•		H.	•	3.1	<u>.</u>
Mean			5.9	6.2	•	•	6.5		30.2	13.5	
LSD (.05)			6.0	6.0	8.0	6.0	9.0	•	4.6	•	4
C.V. (%)			9.5	8.6	7.3	•	6.1	12.1	9.5	40.6	0
F value			20.4**1	16.4**1	2	9.4**	**25.5**	1.5NS	2.0**	40.3**	17.7**

ERWINIA/POWDERY MILDEW EVALUATION OF MONOGERM POPULATIONS AND LINES, SALINAS, CA., 1999 TEST 3699.

40 entries 1-row plot	40 entries x 3 reps., sequential 1-row plots, 17 1/2 ft. long						S S	Planted: A Inoc. Ecb: Scored Ecb:	April 13, July 14, D: Nov. 12	1999 1999 7, 1999
Varietv	Description	Ã	Powdery	Mildew	Score	•	Stand	Harvest Count	Erwinia	Rating
		08/23	08/31	80/60	10/04	Mean	No.	No.	Id	8H
Monogerm F	Monogerm populations									
7835	6833, %, 6834%aa x A	7.0	7.0	7.7	7.7	7.3	30.3	33.3	16.8	74.0
8835m	7835, mmaa x A	6.7	6.7	•	•			31.3	15.0	75.4
8835HO	7835H50 x 7835	•	•	7.3	7.3	6.9	30.7	32.0	21.6	65.9
8835H50	C790-15CMS x 7835	6.7	6.7	7.3	•	6.9	29.0	31.0	•	57.9
7838	6828,6836, aa x A	6.3	6.3	•	•	6.7	26.7	Ξ.		
8838m	7838mmaa x A	•		•			28.3	9.	20.2	6
8838HO (B)	7838H50 x 7838	5.7	5.7	6.3	0.9	5.9	32.0	33.0	9.5	81.5
8838H50	C790-15CMS x 7838	6.3	6.7	•	•		32.7		14.5	H
US H11	Ecb resist. ck.	7.7	7.3	8.3				33.3		89.7
E740	Inc. E840, Ecb susc. ck.	•	0.6	0.6	8.0	•	30.3	31.0	77.2	
6869m	5869mmaa x A	6.7		7.3	•	7.3		ë.	20.3	68.5
7869NB	NB-RZM 5869	6.7	6.7	•	7.7		32.0	33.7	•	67.3
8869m	RZM 7869-#(C)	6.0			_	•	28.0	28.7	ω.	•
0Н6988	7869HO x RZM 7869-#(C)	7.0	7.0	8.0	8.3	7.6	30.7	Ξ.	29.5	0
m0688	RZM 7890, RZM-%S 6890, 5890	6.7	6.7	7.0	7.7	7.0	28.0	28.7	4.9	85.1
8810M	RZM 7810NB	6.3	6.7	7.3	7.3	6.9	29.3	29.7	31.5	51.1
8848M	RZM 7848M	7.0	•	7.3	•	•	31.7	\vdash		
8833	RZM, T-O 7833-#(C), 7834-#(C)	•	•	•	8.3	•	30.3	0	19.8	70.3
8836	T-O 7836-#, 7837-#	7.0	7.0	8.0	7.0	7.3	30.7	31.7		
97-546	Inc. F82-546 (C546)	7.3	7.3	7.3	8.0	•	24.7	9	3.4	91.4

(cont.)

Varietv	Description	Ро	Powdery	Mildew	Score		Stand	Harvest	Erwinia 1	Rating
		08/23		80/60	10/04	Mean	No.	NO.	DI	H%
Monogerm li	lines									
8829-3	Inc. 5829-3 (A,aa), C829-3	7.0	•		8.0	•	œ.	Н	m	9
8829-3H50	x 5829-3	6.7	•	•	7.7	•	7.	∞	•	ė.
8831-3	3831-	6.3	5.7	7.0	6.7	6.4	28.7	29.3	48.6	39.9
8831-3H50	x 5831-3	6.7	•	•	7.0	•	œ.	0	•	7 .
8831-4	T-O 7831-4-#, C831-4			•	•	•	œ	σ.	45.7	ю
8831-4HOM	Ω,	5.7	6.3	7.0	6.7	6.4	24.0	25.3	•	76.9
8833-5	833	•		•	•	•	ري د	9	9.9	a.
8833-5H50	k 5833-5, C8	•		•	•	•	9	7.	•	66.4
8833-12	Inc. 5833-12 (A,aa), C833-12	7		8.7	•		4.	ø.	<u>ი</u>	•
8833-12H50	3 x 5833-12, CE	9		7.7	•		ω ω	œ	33.5	42.4
E740	Inc. E840	0.6	0.6	0.6	8.0	8.8	29.0	30.7	9	•
US H11	Ecb resist. ck.			7.3	•	•	ω.	o.	5.0	83.1
Topcross hy	hybrids with monogerm lines									
X869H5	× ¥769	4.3					7.	9	•	δ.
Y869H27	×	4.7	5.0	5.7	6.7	5.5	27.7	30.0	15.8	70.3
X869H29	×	7.0					7	60	•	ö
Y869H46	7869-6HO x Y769	0.9						Ή.	•	ė.
X869H45	×	•	•	0.	7.0		σ.	8	Ξ.	
Y869H35m	7835mmaa x Y769	•	•	0.	7.0		ö	2	ω̈́	71.0
X869H38m	7838mmaa x Y769	6.0	5.7	6.3	6.7	6.2	32.0	34.3	11.6	75.8
х869н69	7869aa x Y769	•	•	•	•		o.	m	i.	78.4
Mean		9.9	•	•	•	•	29.0	30.4	Η.	9
LSD (.05)		1.1	6.0	6.0	0.8	0.7	4.6	4.3	14.0	20.8
C.V. (%)		10.6	•	•	•	•	9.7	8.8	о	9.
F value		4.8*	•	5.4**	•	ω.	•	•	w.	•

CERCOSPORA LEAF SPOT EVALUATION OF SALINAS ENTRIES, 1999

Ft.

		Ft.				
Variety	Description	Collins	Shak	opee	Ita	ly
		Sep 22	RR	Mean	07/27	08/17
					<u> </u>	30/2/
97-SP22-0	LSR-AR check	3.3	1.5	4.2	1	6
B4430R	L4330.8052, 3-10-97 (CS check)	7.3	2.3	5.7	6	8
	Resist. check	4.3			1	5
Monodoro			2.8	3.6	1	5
Ippolita	Resist. check	4.7	3.1	3.5		
Rifle	Commercial check	6.5	3.1	4.9		
Y869	RZM Y769, C69	5.0	3.2	4.0		
Y875	RZM Y775	5.5	1.9	4.7		
CR811	RZM CR711, CR09/10	4.7	2.6	3.9	3	7
CR812	RZM CR712	5.3	2.9	4.1	4	7
CR813	RZM CR713	5.5	3.3	3.6	3	7
01.010			0.0	• • •	J	, i
8932MCT	7932CT, x A	6.3	3.0	4.4		
EL-02	RZM EL (Rz x sm.root)	4.7	2.4	4.5		
EL-04	RZM EL (Rz x sm.root)	5.0	2.6	4.4	_	
R827 (C27)	RZM R727A,B	5.3	2.5	4.7	3	7
R726 (C26)	RZM-ER R526, C26	5.8	2.8	4.6	3	6
US H11	LSS check	4.8	3.2	4.4		
Dorotea	Resist. check		3.3	3.7		
B4776R	Commercial check		3.3	5.4		
8835	7835aa x A		3.3	4.7		
8935	Inc. R776-89-5H13		3.0	4.2	3	7
0,55	1116. 11770 05 31113		3.0	4.2	3	,
8931	RZM 7931aa x A				2	7
					3	7
CR711	RZM CR11 (C) aa x A				3	7
CR712	6931aa x CR11(C)				3	7
R709-1	CR-RZM R509-1				3	7
C76-89-5	Inc. C76-89-5				3	7
Gabriela	Susc. check				6	8
LSD (.05)		1.0	1.2	0.7		
SP351069.0	LSS check	6.5				
(FC504 x FC502/		3.3				
(10304 X 10302)	27 % 5122 0	3.3				
From CBGA Coded	Test					
Beta 4430R						
	Commercial hybrid			5.6		
US H11	Susc. check			4.2		
Mod. resist. ch				3.3		
Mod. susc. chec	ek			4.8		
Susc. check				5.3		
Resist. check				2.8		

Ft. Collins: Test by L. Panella, USDA-ARS
Shakopee: Test by Betaseed run by M. Rekoske and J. Miller
Italy: Test by E. Biancardi, Rovigo, Italy

Test by E. Biancardi, Rovigo, Italy Italy:

RR = root rot score (Aphanomyces)

Planted: November 3, 1998 Not planted for havest

160 entries x 3 reps., sequential 1-row plots, 17.5 ft. long

		Stand	Emergence				%Downy
Variety	Description	Count	Score	GC.	% Bolting	1	Mildew
		Mean	1/21	07/28	08/26	10/06	05/26
Checks US H11	113102	26.0	4.0	14.0	ω	20.0	
SS-NB3	Spreckels, 1996	80	•	•	22.2	32.2	35.9
97-C37	Inc. U86-37	<u>ი</u>	•	•	•	0	o.
U86-37		Ή.	•	•	8	8	<u>ი</u>
97-SP22-0			4.7	1.	8	8	
97-US22/3	Inc. Y009 (US22/3)	ω.	•	8	ω.	9	ω.
97-US75	Inc. 268 (US75)	•	5.0	19.5	•	•	9
B4776R	Betaseed	28.0	•	2.	9	·	
Multigerm, op	open-pollinated lines						
12	RZM R778%, (C78)	7.	4.3	8	т М	თ	9
R778% (Iso)	R578, R578/2	28.3	4.3	18.9	32.0	28.4	59.9
R778 (Iso)	RZM-ER R578, R578/2, R578%	0	٠	7.	8	ω.	8
R878 (Sp)	Inc. R778, R778%	8	3.3	7.	œ ·	Η.	4.
R880	RZM R780, (C80)	ω	4.7	2	ъ.	9	Ή.
R882 (Sp)	1 - 4	œ	•	•	•	34.3	4.
R881 (Iso)	RZM R776, R781, R681, (C82)	28.3	5.0	38.7	37.5		42.6
R776	RZM-ER R576 (C31Rz)	œ	•	•	•	9.1	
R781	RZM-ER R581	σ.	•	5.	ο.	•	0
R770	RZM-ER R570	œ	5.0	о О	7.	ω.	2
R879	RZM R779, (C79-1,Rz)	27.7	4.0	35.8	33.5	35.8	38.0
R736	RZM R636, (C79-8,R22)	9	4.3	Η.	Ή.	o,	7 .
R836	RZM R736, R743 (C79-8,R22)	0	•	ъ.	9	4.	9
R753	RZM R653	თ	•	8	Ή.	9	
R853	RZM-ER-%S R653	29.3	5.0	23.1	28.7	27.2	33.0
R854	RZM R754	7.	•	ω.	ω.	2.	

EVALUATION OF BREEDING LINES AND POPULATIONS FOR NONBOLTING, SALINAS, CA., 1998-99 TEST 299.

Description
Smooth)
Smooth)

TEST 299. EVALUATION OF BREEDING LINES AND POPULATIONS FOR NONBOLTING, SALINAS, CA., 1998-99

Mildew *Downy 05/26 45.9 39.2 25.8 43.8 32.2 33.3 34.9 49.4 80.3 37.9 47.2 32.7 23.0 9.0 27.4 42.3 14.4 10/06 42.5 26.4 46.9 40.9 65.5 56.0 6.09 33.8 47.5 39.2 43.8 33.0 39.3 43.7 41.5 7.7 58.4 35.8 9.09 37.2 38.5 38.9 % Bolting 08/26 34.4 29.9 40.0 54.2 23.4 25.3 45.7 36.9 65.2 58.4 60.09 57.5 35.9 27.7 55.4 60.5 45.1 56.1 07/28 24.2 30.8 28.2 20.7 43.2 27.3 32.0 35.3 58.9 52.9 54.7 60.4 28.9 2.8 55.0 27.6 43.9 37.1 16.4 30.3 35.9 32.1 32.1 31.1 Emergence Score 1/21 5.0 4.7 5.0 4.3 5.0 5.0 4.7 4.7 4.7 4.3 4.7 5.0 5.0 4.3 4.7 4.7 27.0 27.7 28.0 27.3 25.0 29.7 Stand Mean 29.0 28.7 28.3 28.3 29.0 27.0 27.3 28.0 28.0 27.0 28.0 27.7 27.0 27.0 26.7 27.7 27.3 27.7 Count (CP02) (CP01) XZM 7931aa x A, (popn-931) RZM-PMR 6203-#,6208-#(C) RZM-PMR 6211-#,6217-#(C) CR-RZM R509-#, R510-#(C) 6205-#,6206-#(C), 6201-#,6202-#(C), Description Inc. N629,N630 (galls) RZM 7924, ... aa x 924(C) Sf, Aa populations & lines $Z625-\#(C)aa \times Z31(C)$ $Z630-\#(C)aa \times Z31(C)$ 3ZM Z731, Z730aa x A 7932CT,7201, aa x A Inc. 7932CT,7201...A RZM CR711 (CR09/10) 7931aa x RZM 7926 7204-7216CMS x A CR-RZM R509A-10 5931aa x 931(C) RZM 7926 (A, aa) CR-RZM R509A-9 RZM 7926aa x A CR-RZM R509A-1 Inc. 6260-#,... 6869mmaa x A RZM CR713 RZM CR712 Inc. Inc. Multigerm, Variety 8926 (Iso) 8926 (Sp) 8932HO (M) 8932H69 R710-10 8932Am R709-9 7932CT R709-1 CR811 **CR812 CR813** R710 N730 8932 2730 P811 P812 P813 P814 8924 2725 7931 8931 2831

EVALUATION OF BREEDING LINES AND POPULATIONS FOR NONBOLTING, SALINAS, CA., 1998-99 TEST 299.

		(cont.)					
		Stand	Emergence				%Downy
Variety	Description	Count	Score 1/21	07/28	% Bolting 08/26	10/06	Mildew 05/26
Multigerm, S ^f	Sf, Aa populations & lines (cont.)						
(as)	Inc. R776-89-5H13Aa			7.		ري	5.
Iso)	~	27.3	4.7	19.5	23.3	28.0	35.4
	RZM R776-89-5H31	ω.	•	8		9	0
	RZM R776-89-5H11	6	•	Ŋ.		80	ij.
	RZM Z731H11	80		2	4	80	Η.
	RZM Y769H31	8	•	رى	9	4.	8
х869н31	7931aa x Y769	28.7	5.0	29.1	43.0	44.2	17.4
	Inc. 6264-#(C)	9	•	9.	œ.	4.	ë.
-70	RZM-ER-%S 6913-70, (C913-70)	0		•	2.		
8918-12	RZM-ER-%S 6918-12	7 .	•	8	•	•	o.
_	RZM 7918-21	7.	•	•	H.	л	•
8911-4-10M	RZM-ER-%S 6911-4-10	æ.	•	•	•	•	•
•	Inc. 6925-19	7.	4.0	ς.	•	o.	ö
	Inc. Z625-6 (A,aa)	27.3	4.7	31.7	32.9	29.1	47.6
2825-9		œ	•	4.	•	2	Б.
_	Inc. Z630-11 (A,aa)	9	3.7	ω.	9	9.	4
8929-41	Inc. 6929-41 (A, aa)	ω.	•			•	1.1
2	Inc. 6929-72 (A,aa)	。	•	•	•	•	4.
02	2	28.0	5.0	24.1	27.7	28.9	29.9
12		9	•	。	0	ö	2
8929-114	Inc. 6929-114 (A,aa)	7.	•	34.7	•		4
15		9	4.0	。	41.7	4	•
8929-133	Inc. 6929-133 (A,aa)	25.7	4.3	g.6	0.6	11.7	20.4
53		ω	4.7	•		•	3.4
54	Inc. 6929-154 (A,aa)		4.3	34.0	•	•	•

Mildew 05/26 %Downy 58.7 8.4 49.6 41.2 55.3 10/06 22.3 35.8 20.7 17.6 0.0 57.7 80.8 % Bolting 08/26 54.3 23.5 3.4 07/28 52.1 86.5 0.0 17.5 1.2 20.1 Emergence Score 1/21 4.7 4.0 4.7 28.0 Stand Mean 28.3 Count 27.0 26.0 26.7 26.3 Sf, Aa populations & lines (cont.) Description Monogerm, St. As populations & lines Inc. 6930-102 (A,aa) (A,aa) Inc. 6930-19 (A,aa) (A,aa) (A,aa) (A, aa) (A, aa) 6927-29 68-0869 6927-30 6927-33 6927-37 Inc. Inc. Inc. Inc. Inc. Variety Multigerm, 8930-102 8930-19 8930-39 8927-30 8927-33 8927-29 8927-37

MOHOGETH, 3	Molioderii, S, Aa poputations & IInes						
6546	Inc. F82-546, (C546)				20.6	30.6	12.5
6562	Inc. F82-562, (C562)	26.0	3.3	37.2	20.5	28.5	6.4
6718	Inc. U83-718, (C718)		•		7.4	0.9	7
6762-17	Inc. 0762-17,2762-17, (C762-17)	80	•	-	•		33.2
7835	6833, 6834% 6834% x 835 (C)	_		σ	Ľ	Ľ	Ľ
8835m) 	_	•	, –) α	α	} c
8835M	7835Maa x A	27.3	0.0.0	25.6	29.3	32.9	14.7
8835HO	7835H50 x 7835	9	•	2.	6	6	
8835H50	C790-15CMS x 7835	· 80	•	4.	2	6	
7838	6828,6836,6837, aa x 838 (C)	28.3	5.0	31.8	36.3	40.9	4.
8838m	7838mmaa x A	7.	•	ω.	т М	6	H
8838M	7838Maa x A	7.	•	4.	.	37.0	30.5
8838H11m	5911-4mmaa x 7838	25.0		7.	80		υ.
8838HO (A)	7838H10 x 7838	27.3	•	ω.	80	8	4.
8838HO (B)	7838H50 x 7838	27.7	4.0	16.8	19.2	24.1	ω.
8838H50	C790-15CMS x 7838	27.7	4.3	9	9	2	19.2
7869NB	NB-RZM 5869m(A,aa), (C869)	σ.			9	7.	w.
8869m	RZM 7869-#(C)m	ъ Э		5	4.	<u>ი</u>	ä
8869но	7869HO x RZM 7869-#(C), (C869CMS)	28.0	5.0	21.5	28.4	35.5	10.8
8890m	RZM 7890,6890,5890m (A,aa),(C890-1)	7.		5	4.	4.	e.

EVALUATION OF BREEDING LINES AND POPULATIONS FOR NONBOLTING, SALINAS, CA., 1998-99 TEST 299.

Varietv	Description	Stand	Emergence Score		% Bolting	_	%Downy Mildew
		Mean	1/21	07/28	08/26	10/06	05/26
rm, S ^f		,			,		
0Н0688	7890HO x RZM 7890,, (C890-1CMS)	o	•				
8810m	RZM 7810NBm	25.0	•	13.6	12.2	14.9	21.2
8810M	RZM 7810NM	89	•		•		
8848m	RZM 7848m (A,aa)	7.	4.7	•	•	•	
8848HOm	7848H88m x RZM 7848	ဖ	4.3		34.7	33.6	
8833	7833-#	27.7		58.9	2	8.8	. 6
8833450	15CMS x 7833-#(C	· 00				7 67	α
8836		26.3	4.3	1.3	2.6	m	9.5
ВВЗКНОМ	#-7835#-9836-#-X837-#	7	6	6 7		12.2	16.4
0000	6-0000					ш	
8829-3	(5829-	7.07	4 J. 1	0.0)) (ກ. ເ	
8829-3H50	-15 CMS $\times 5829-3$		•	0.0	•	0.9	17.1
8831-3	Inc. 5831-3 (A,aa), (C831-3)	5	4.0	ж В	•	9. 8	
8831-3H50	C790-15CMS x 5831-3	9.	•	20.9	26.1	25.1	ъ.
8831-4	T-0 7831-4-#(C) (A,aa), (C831-4)	28.0	4.7	0.0	0.0	9.7	19.1
8831-4HO	$8131-4HOM \times 7831-4-\#(C)$	7.	•	1.3	•	2.5	æ.
8833-5	Inc.5833-5 (A,aa), (C833-5)	6	4.3	•	24.2	•	0
8833-5H50	C790-15CMS x 5833-5	7.	4.7		17.8	21.8	7.
8833-12	Inc. 5833-12 (A,aa), (C833-12)	27.3	4.3			41.9	9
8833-12H50	C790-15CMS x 5833-12	•	5.0	41.3	54.3	65.5	13.3
8911-4-7	STO 7911-4-7-#(C), (C911-4-7)	25.3	g.8		44.7	40.1	0
8911-4-7H50	$6911-4-7HO \times 7911-4-7-\#(C)$	26.3	4.0	42.9	39.5	39.0	
8818-1 (C)	Inc. 6818-1mm (A,aa)	4.	2.0	5.6	5.4	8.2	14.1
8818-2 (C)		26.3		•	6.4	6.4	
8818-6(C)		5	3.0	•	4.1	5.4	24.5
8818-11 (C)	Inc. 6818-11mm (A,aa)	21.3	2.0	7.7	6.1	9.2	6.3
8818-12 (C)	Inc. 6818-12mm (A,aa)	Ŋ	4.0	0.9	11.5	18.1	

EVALUATION OF BREEDING LINES AND POPULATIONS FOR NONBOLTING, SALINAS, CA., 1998-99 TEST 299.

(cont.)

1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		Stand	Emergence	d	14.1		%Downy
variety	Description	Mean	1/21	07/28	8 BOLCING 08/26	10/06	05/26
Monogerm, S ^f , 1 8818-21 (C)	Monogerm, S ^f , Aa populations & lines (cont.) 8818-21(C) Inc. 6818-21mm (A,aa)	25.0	2.7	0.0	0.0	0.0	8.9
8818-1B	Inc. 6818B-1	26.7	4.3	26.1	25.0	24.8	13.4
8818-1BHO	C790-15CMS x 6818B-1	28.0	5.0	27.1	42.7	49.5	9.6
8818-2B	Inc. 6818B-2	27.0	4.0	17.4	18.6	31.9	49.5
8818-2BHO	C790-15CMS x 6818B-2	0.7	0.0	g	no plants		
F92-790-15	Inc. 1790-5 (C790-15) (921194)	27.0	э. Э.	38.3	48.2	45.5	10.0
Mean		27.3	4.4	27.4	33.1	34.1	31.8
LSD (.05)		2.7	0.8	17.3	17.3	18.5	19.8
C.V. (%)		6.1	11.1	39.3	32.6	33.7	38.8
F value		**0.6	8.0**	8.7**	10.1**	7.3**	6.7**

accurately make. Due to severity of bolting and diseases, the second bolting counts (8/26/99) are probably the winter-spring of 1999 was colder than normal. Much higher levels of bolting were experienced than for In addition, a number of years. Downy mildew (Peronospora farinosa) appeared in early spring and by mid-summer became Sclerotium rolfsii (southern root rot) also became severe in Downy mildew affected plant growth and survival and probably rate and percent bolting. Counts this planting. Due to plant death and rotting, bolting counts later in the season were difficult to NOTES: Bolting tests were planted earlier than in previous years to get greater induction. the most accurate and show the best differential levels between entries. were made based upon obvious top symptoms. severe.

TEST 199. EVALUATION OF TESTCROSS HYBRIDS FOR NONBOLTING, SALINAS, 1998-99

Planted: November 3, 1998 Not harvested for yield 80 entries x 3 reps., sequential 1-row plots, 17.5 ft. long

Varietv	Description	Stand Count	Emergence Score	•	% Bolting		%Downy Mildew
		Mean	1/21	07/28	08/26	10/06	05/26
US H11	113102	28.3	4.7	13.2	6	5	18.6
SS-NB3	Spreckels, 1996	Η.	5.0	9	24.3	24.2	4
B4776R	Betaseed 4776R.7653 (3-27-98)	30.7	5.0	31.6	46.9	m.	26.2
B4035R	Betaseed 4035R (7-10-97)	30.0	5.0	40.0	52.3	51.2	27.0
Rizor	Holly HH108, 9-3-97	31.0	5.0	60.2	73.7	58.3	7.
Rifle	Holly, 9-16-98	29.3	•	49.0	æ	4.	1.
SS-778R	Spreckels, X782402	•	5.0	38.5	•	57.8	33.6
5KJ0142	Betaseed (8-18-97)	28.3	4.7	30.5	49.3	44.6	•
R878%H50 (Iso)	C790-15CMS x RZM R778%		5.0	m.	45.0	46.2	13.4
R878H50 (Sp)	C790-15CMS x R778, %	6	•	7.	ო	80	6
R878H69	7869aa x R778,8	28.3	•	29.5	31.6	34.0	13.9
R878H55	7835H50 x R778,%	6	3.7	4.	رى	7.	9
R878H58	7838H50 x R778,%		3.7	24.6	34.7	35.7	24.0
R876-89-5NBH50	C790-15CMS x RZM-%S R576-89-5NB	•	4.0	19.9	9	27.1	9.7
R876-89-5H50	C790-15CMS x RZM-%S R576-89-5	œ	5.0	•	32.5	•	8.1
X882H50	C790-15CMS x R781,R776,	28.7	4.7	9		Ή.	19.0
Y882H37	4807HO x R781,R776,	28.7	5.0	19.7	39.4	43.9	11.7
X882H27	6831-4HO x R781,R776,	27.0	•	21.3	30.0	7.	H.
Y882H38m	7838mmaa x R781,R776,	9	4.7	30.1	0	51.2	21.3
Y875H50 (Iso)	C790-15CMS x RZM Y775	29.0	5.0	œ.	35.6	4	m.
Y875H50 (Sp)	C790-15CMS x RZM Y775,	28.7	4.7	17.4	23.2	24.4	2
X875H37	4807HO x RZM Y775,		•	ю Э	9	53.1	m
X875H27	6831-4HO x RZM Y775,	•	4.7	13.1	24.9	22.6	16.8
Y868 H50	C790-15CMS x RZM Y678	30.3	5.0	•	4.	23.0	m.

(cont.)

Varietv	Description	Stand Count	Emergence Score	%	Bolting		%Downy Mildew
		Mean	1/21	07/28	08/26	10/06	05/26
Y866H50	C790-15CMS x RZM Y766	29.7	5.0	26.1	43.1	46.7	28.3
X867H50	C790-15CMS x RZM Y767 (C67)	29.7	5.0	5	45.7		Б.
X871H50	C790-15CMS x RZM Y771	9.	•	26.2	7	48.2	24.1
Y872H50	C790-15CMS x RZM-%S Y672	29.3	5.0	e.	30.6	36.2	o.
Y872H50	C790-15CMS x RZM Y772 (C72)	80	4.7	⊢.		52.4	22.2
Y869H50	C790-15CMS x Y769		4.7	0	8	41.9	18.0
R879H50	x RZM	28.7	5.0	28.0	•	36.1	18.6
R836H50	C790-15CMS x RZM R736		5.0	2	51.1		19.3
R854H50	C790-15CMS x RZM R754	ω.	5.0	32.1	41.7		13.2
R873BH50	C790-15CMS x RZM Y773	30.3	5.0	31.1	•	38.6	20.9
R835H50	C790-15CMS x RZM R735	6	•	•	45.7	H.	
8931H50	C790-15CMS x RZM 7931	œ.	5.0	11.9	•		о
8931H46	7869-6HO x RZM 7931	8	4.3	1.	7	5.	9
8931H38m	7838mmaa x RZM 7931	œ.	•	2	7	ω.	25.7
8924H50	C790-15CMS x RZM 7924	28.3	5.0	23.7	37.8	33.0	5.
Z831H50	C790-15CMS x RZM Z730,Z731	o.	•	8	2	7 .	
Z831H37	4807HO x RZM Z730, Z731		4.3	19.7	32.3	0	0
CR812H50	C790-15CMS x RZM CR712	28.7	5.0	47.7	60.3	6	25.7
CR813H50	C790-15CMS x RZM CR713		•			58.9	급.
8926H50 (Iso)	C790-15CMS x RZM 7926	ω.	5.0	23.7	41.9	5.	18.6
8926H50 (Sp)	C790-15CMS x RZM 7926	7.	5.0	0	•	ი	7.
8926н37	4807HO x RZM 7926	•	4.7	31.3	46.7	40.2	37.7
8932H50	C790-15CMS x 7932CT,7201		•	34.3	8	7.	19.9
8932H38m	7838mmaa x 7932CT,7201	29.7	4.7	43.9	50.7	48.4	0

EVALUATION OF TESTCROSS HYBRIDS FOR NONBOLTING, SALINAS, 1998-99 TEST 199.

	%Downy Mildew 05/26	20.0 31.0 34.8 21.1	16.7 16.1 34.4 29.9	31.6 18.1 17.5 4.6	23.2 11.5 39.7 14.5	19.0 24.4 19.2 3.5 13.8 25.7 25.7
	10/06	36.3 25.6 40.8 33.2	25.9 28.4 29.4 51.4	28.7 56.8 44.7 18.5	29.2 48.9 3.4 57.0	41.6 45.6 38.6 20.0 4.6 31.0 16.0
	Bolting 08/26	33.0 26.6 33.2 40.5	32.8 26.1 30.5 51.7	33.1 50.0 38.0 16.3	28.1 48.8 1.1 54.7	4 4 5 0 0 6 3 3 3 4 4 1 8 0 8 8 1 1 8 0 5 1 1 8 0 5 1 1 1 7 0 8 1 1 7 0 8 1 1 7 0 8 1 1 7 0 8 1 1 7 0 8 1 1 7 0 1 1 8 1 1 7 0 1 1 8 1 1 7 0 1 1 8 1 1 7 0 1 1 8 1 1 7 0 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
	8 07/28	16.1 13.8 16.5 24.8	15.3 20.5 21.0 32.6	19.6 22.7 27.8 6.9	12.3 25.6 1.1 42.5	36.0 23.3 24.9 11.7 16.9 10.0
	Emergence Score 1/21	4 5 5 5 4 5 0 0 0 0 0 0 0	5.0 5.0 5.0	5.0 4.7 7.0	 	
(cont.)	Stand Count Mean	29.3 28.7 28.0 28.0	28.3 28.3 28.3	29.3 28.0 28.7 29.0	30.0 29.7 29.3 30.0	29.7 30.0 30.0 28.3 29.3 29.0
	Description	C790-15CMS x RZM R776-89-4H13 C790-15CMS x R776-89-5H13 4807HO x R776-89-5H13 7838mmaa x R776-89-5H13	C790-15CMS x RZM R776-89-5H31 C790-15CMS x RZM R776-89-5H11 C790-15CMS x RZM Z731H11 C790-15CMS x RZM Y769H31	C790-15CMS x RZM-ER-%S 6913-70 C790-15CMS x RZM-ER-%S 6918-12 C790-15CMS x RZM 7918-21 C790-15CMS x RZM-ER-%S 6911-4-10	C790-15CMS x 6925-19 C790-15CMS x 6929-41 C790-15CMS x 6929-72 C790-15CMS x 6929-102	C790-15CMS x 6929-112 C790-15CMS x 6929-114 C790-15CMS x 6929-115 C790-15CMS x 6929-133 C790-15CMS x 6929-153 C790-15CMS x 6929-154 C790-15CMS x 6930-19 C790-15CMS x 6930-39
	Variety	8935H50 (Iso) 8935H50 (Sp) 8935H37 8935H38m	8936H50 8937H50 8938H50 8939H50	8913-70H50 8918-12H50 8918-21H50 8911-4-10H50	8925-19H50 8929-41H50 8929-72H50 8929-102H50	8929-112H50 8929-114H50 8929-115H50 8929-133H50 8929-153H50 8929-154H50 8930-19H50

27.8 43.8 33.3 25.1

8.9 49.5 40.5 53.2

6.7 43.9 39.3 52.1

6.7 27.0 25.0 46.4

5.00

30.0 29.7 28.0 29.3

C790-15CMS x 6930-102 C790-15CMS x 6927-29

8930-102H50

8927-29H50 8927-30H50 8927-33H50

C790-15CMS x 6927-30 C790-15CMS x 6927-33

EVALUATION OF TESTCROSS HYBRIDS FOR NONBOLTING, SALINAS, 1998-99 TEST 199.

(cont.)

		Stand	Emergence				%Downy
Variety	Description	Count	Score	olo .	Bolting		Mildew
		Mean	1/21	07/28	<u>3 08/26 1</u>	10/06	05/26
8927-37H50	C790-15CMS x 6927-37	28.3	5.0	61.4	76.7	71.8	26.0
Z825-6H50	C790-15CMS x Z625-6	29.3	4.7	30.7	51.0	57.8	28.5
Z825-9H50	C790-15CMS x Z625-9	29.0	5.0	28.7	52.9	50.5	32.1
Z830-11H50	C790-15CMS x Z630-11	29.0	5.0	41.5	56.3	49.5	49.6
Mean		28.9	4.8	25.7	38.5	38.4	23.7
LSD (.05)		2.3	0.5	14.2	15.5	16.5	18.9
C.V. (%)		5.0	6.3	34.2	25.0	26.6	49.5
F value		1.2NS	3.6**	5.3**	6.5**	ນ. 3**	1.9**

Notes: See notes for Test 299. In general for % bolting and % downy mildew infection, there is good correspondence between the line (Test 199) and its hybrid. Particularly for downy mildew, lines with high resistance produced hybrids with moderate resistance. For some lines and hybrids, the relationship for bolting was less clear cut.

EVALUATION OF TOPCROSS HYBRIDS FOR NONBOLTING, SALINAS, 1998-99 TEST 999.

r 3, 1998 _Y ield	%Downy Mildew 05/26	7.8		•	~ თ ი			15.6	0.0	14.9	Ή.	8.3	•	4	•	7	12.3	•	•		34.2	0.6	•	4.	15.6	。
embe; for	10/06	Ι .			37.5 38.55			œ.	38.5	œ ·	7.	9	Б.	-	0	4.	39.4	ω.	4		ω.	6	9	4	15.3	9
Planted: Nov Not harvested	Bolting 08/26	-	6	0 1	30.5	4		9	38.5	о О	ω.	9	e.	4	ω.	2	32.9	9	9		23.5	61.7	4.	•	8.5	•
ωZ	8 07/28			<u>.</u>	25.6			2	28.4	о О	5	6	4.		2	о О	31.5	6	ω.		14.9	•	•	•	8.6	•
	Emergence Score 1/21	Ι .	4.7	•	4.0	•		•	3.7	•	4.7	4.0	•	4.3	4.7	4.3	4.7	•	4.3		4.3		•	•	4.0	•
	Stand Count Mean	17.0	7.	ت	16.7	•		5.	14.3	9	5	14.7	5.	16.3	•	15.7	16.0	9	16.7		16.0	•	5	ς.	15.3	9
reps., sequential 1 ft. long	Description	Spreckels, 1996			Inc. 1/69 (C69) 4807HO (C306/2) x Y769	69	rids	7835mmaa x Y769	7838mmaa x Y769	7869mmaa x Y769	7931aa x Y769	7932CTMaa x Y769	7204-7216CMS x Y769	7890HO x Y769	7869HO x Y769	7835H50m x Y769	7838H50m x Y769	7838H10m x Y769	7848H88m x Y769	pl	C833- 5aa x Y769	C833-12aa x Y769	C829-3aa x Y769	$C831-3aa \times Y769$		$C911-4-7HO \times Y769$
96 entries x 3 1 1-row plots, 11	Variety	Checks SS-NB3	US H11		Y869 (Sp) Y869H37	X869H50	Population Hybrids	Y869H35m	Y869H38m	х869н69	Y869H31	х869н30М	х869н32	х869н88	X869H70	X869H55m	X869H58m	X869H59m	Y869H49m	Testcross Hybrids	X869H5	Y869H12	X869H29	X869H4	X869H27	X869H7

(cont.)

		Stand	Emergence				%Downy
Variety	Description	Count	Score	8	% Bolting	J	Mildew
		Mean	1/21	07/28	08/26	10/06	05/26
Testcross Hybrids	s (cont.)						
Į.	_7869-6HO x Y769	15.7	3.7	23.2	34.6	44.6	22.8
X869H45	C867-1HO x Y769	9	•	46.9	т М	63.5	34.8
X869H17	7817HO x Y769	15.0	3.7	17.8	17.8	15.6	6.7
X869H18	7818HO x Y769	9	4.7	23.8	7.	•	20.1
X869H19	7818H50 x Y769	9	4.3	8	8	0	•
х 869H20	7818-4H50 x Y769	15.7	4.0	e.	e.	32.3	21.7
Y869H21	7818-14H50 x Y769	9	4.7	21.4	8	44.3	15.3
X869H22	7818-22H50 x Y769	15.3	•	17.2	17.2	7.	
Ү 869H23	7818-23H50 x Y769	5	5.0	25.6	9	33.6	16.9
Topcross hybrids	onto 818-#s						
X869H15-1B	6818-1Baa x Y769	ъ.	4.3	4.	7.	ω.	4.8
Y869H15-2B	6818-2Baa x Y769	16.0	4.3	14.6	20.4	16.5	5.9
Y869H15-1	6818-1aa x Y769	9	4.0	16.2	0	4.	18.5
X869H15-2	6818-2aa x Y769	15.3	4.0	•	18.8	25.3	•
X869H15-6	6818-6aa x Y769	16.0	4.3	6.3	4.2	6.3	18.6
Y869H15-21	6818-21aa x Y769	e.	3.0	•	•	14.8	ij.
Topcross hybrids	onto 808-#s						
У869Н9 - 1	7808- 1aa x Y769	14.0	•	5	45.0	50.0	4.6
1 2	7808- 2aa x Y769	13.7	n. n.	33.7	61.7	61.2	14.3
e I	7808- 3aa x Y769	4	3.7	÷.	19.3	•	0.0
- 4	7808- 4aa x Y769	ъ.	4.0	17.4	21.3	ъ.	16.0
L -	7808- 7aa x Y769	15.3	4.0	6.5		15.1	
& 1	7808- 8aa x Y769	Б.	4.0	10.4	4.	•	0.0
б	7808- 9aa x Y769	15.7	4.0	18.5	37.7	35.8	16.4

EVALUATION OF TOPCROSS HYBRIDS FOR NONBOLTING, SALINAS, 1998-99 TEST 999.

%Downy Mildew	05/26		13.1	3.9	•		20.1	4.0		8	•		•	12.5	2	•	5.		0	o.	•	2	•		•	e.	23.4	ο.	ij.
	10/06		i.	31.3	ت			59.7	•	5	6	5	7	45.3	9	9.	o.		4.	ij.	8	30.7	9		ς.	ω.	33.7	7.	60.4
Bolting	08/26		5	30.6	9		e.	59.7	7.	6	9	7.	5.	49.6	0	о О	4.		e.	Ϊ.	2	32.6	4		œ.	9	29.0	8	5
ογo	07/28		•	18.3	•		0	39.1	5	æ	7.	2.	7.	22.7	ø.	5.	æ.		5.	4.	2	26.7	ი		رى	ж	31.5	9	•
Emergence Score	1/21		4.7	4.0	•		4.0	4.3		3.7	•	4.0	4.3	4.7	4.3	•	5.0		4.7	4.0	•	4.3	•		4.3	•	3.7	•	•
Stand	Mean		5.	15.0	4		ė.	16.3	4.	4.	5.	5	9	16.3	5	4.	7.		ъ.	e.	9	17.0	5		ė.	5	15.0	ø.	4
Description		ls onto 808-#s (cont.)	7808-12aa x Y769	7808-13aa x Y769	7808-16aa x Y769	ls onto popn-869-#s	7869- 1aa x Y769	7869- 2aa x Y769	7869- 4aa x Y769	7869- 5aa x Y769	6	7869- 7aa x Y769	7869-13aa x Y769	7869-19aa x Y769	7869-20aa x Y769	7869-20Baa x Y769	7869-24aa x Y769	lsonto popn-833-#s	7833- laa x Y769	7833- 3aa x Y769	7833-10aa x Y769	7833-11aa x Y769	7833-12aa x Y769	onto popn-83	$7834 - 1aa \times Y769$	7834- 2aa x Y769	7834- 3aa x Y769	7834- 5aa x Y769	7834- 8aa x Y769
Variety		Topcross hybrids	-12	-13	-16	Topcross hybrids	<u> т</u> 869н69 - 1	- 2	4 -	ا 5	9	- 7	-13	-19	-20	-20B	-24	Topcross hybridsonto	<u> Y869Н33 - 1</u>	г 1	-10	-11	-12	Topcross hybrids	X869H34 - 1	- 2	n ۱	I N	ω Ι

TEST 999. EVALUATION OF TOPCROSS HYBRIDS FOR NONBOLTING, SALINAS, 1998-99

(cont.)

Variety	Description	Stand Count	Emergence Score	о. С	% Bolting	ħ	%Downy Mildew
		Mean	1/21	07/28	08/26	10/06	05/26
Topcross hybrids Y869H28 - 9	onto popn-828-#s 7828- 9aa x Y769	15.7	4.3	8	ω.	5.	
-10	7828-10aa x Y769	17.0	4.3	13.7	21.6	23.5	3.9
	onto popn-831-	ı		((
	laa x Y76	5	٠	m ·	•	5	•
1 2	- 2aa x Y76	ك	•	9	Η.	i N	Η.
L - 7	- 7aa x Y76	4.	•	H.	•	•	•
ω Ι	7831-4- 8aa x Y769	15.7	4.0	19.2	23.2	25.5	23.3
ი I	7831-4- 9aa x Y869	4.	•	ö	•	•	•
-10	7831-4-10aa x Y769	4.	•	m.	æ.	40.2	•
Topcross hybrids	onto popn-836-#s						
хв69н36 - 3	7836- 3aa x Y769	•	•		•	5.	
-10	7836-10aa x Y769	15.3	3.7	35.1	43.8	55.1	2.1
-11	7836-11aa x Y769	•	•	2		Б.	9
-14	7836-14aa x Y769	e.	•	2	9.	6	•
Topcross hybrids	onto popn-837-#s						
X869H77 - 1	7837- 1aa x Y769	13.7	3.0	о	7.	19.8	
- 1B	7837- 1Baa x Y769		•	27.9	25.4	•	14.0
1 2	7837- 2aa x Y769	9	•	ω.	Ŋ.	•	•
e 1	7837- 3aa x Y769	16.0	5.0	37.5	54.2	56.3	4.2
- 4	7837- 4aa xY769	4.	•	4.	Ŋ.	o.	•
Topcross hybrids	onto popn-839-#	v		~	~		
	3 6 6		•) () c		i c
7 (200 X	7.01	4, 4 O U	4. Z. Z. C.	0.00	40.1 1.010	N C
	א ממט .		•	ດ ເ	٠, رح	٠ د	•
7 -	7839- 4aa x Y769		•	7.	H	o.	

TEST 999. EVALUATION OF TOPCROSS HYBRIDS FOR NONBOLTING, SALINAS, 1998-99

(cont.)

Variety	Description	Stand	Emergence Score	o∤o	% Bolting		%Downy Mildew
		Mean	1/21	07/28	08/26	10/06	05/26
Topcross hybrida Y869H79 - 5	Topcross hybrids onto popn-839-# (cont.)	16.0	4.3	ر د د	24.2	بر د	α
- 5B	7839- 5Baa x Y769	13.3	3.7	50.4	52.0	60.4	7.6
9 -	7839- 6aa x Y769	15.3	3.7	26.3	26.3	30.7	14.6
-10	7839-10aa x Y769	15.3	4.3	26.3	30.6	26.3	11.0
B4776R	Betaseed, 3-27-98	15.0	3.7	40.7	55.9	61.2	2.2
Mean		15.4	4.0	27.1	36.0	38.6	12.1
LSD (.05)		1.9	0.8	18.0	22.7	23.2	19.1
C.V. (%)		7.6	12.8	41.3	39.2	37.4	7.76
F value		2.4**	3.2**	3.5**	3.7**	3.2**	1.2NS

NOTES: See notes for tests 199 and 299.

(TESTS 199, 299, 2699, 3599, 5899, B399 and B899)

Test 3599 (PM-ERR) ERR-DI 17.8 18.6 5.6 3.6 23.8 7.6 0.1 æ| Powdery Mildew 7.6 9.9 4.7 9.9 6.1 5.5 œ Test 299 (NB-Line) 34.3 3.6 46.0 50.3 70.4 8.4 47.6 15.2 64.3 58.7 30.3 41.2 54.5 54.9 M æ | Bolting 2.4 36.2 9.6 27.8 58.1 3.4 13.6 31.0 **₩**| Test 199 (NB-Hyb) 4.6 32.1 25.7 21.8 26.2 19.7 23.2 31.6 18.1 17.5 28.5 39.3 25.7 13.3 27.8 19.0 8.1 ద ₩| Bolting 60.3 56.4 46.9 51.0 52.9 58.0 33.1 16.3 50.0 18.2 17.8 32.5 83.9 84.9 85.8 83.9 85.3 81.5 85.7 83.6 84.1 85.1 83.2 84.6 84.5 83.1 84.7 RJAP 85.1 Test 2699 (yield) Experimental hybrids with S₁ pollinators S₁ pollinators from MM, VY, S^f, Aa, Rz popns R878H50 13129 16.41 Sucrose 15.86 15.96 16.49 16.79 16.10 16.23 16.30 16.64 17.13 17.29 18.02 16.24 16.24 16.74 15.96 18.21 16.96 16.16 16.81 16.01 16.40 16.41 Sugar Yield 14706 14356 14773 14840 14342 15044 13510 14374 13288 15069 14377 14107 14106 13953 14145 14709 14182 14744 1bs 14662 12922 14911 R876-89-5H50 8911-4-10H50 8930-102H50 Variety 8925-19H50 8913-70H50 8918-12H50 8918-21H50 Z830-11H50 8930-19H50 8930-39H50 Z825-6H50 Z825-9H50 R709-1H50 R709-9H50 CR812H50 CR813H50 R882H50 SS-432R 8931H50 B4776R Rifle

EVALUATION OF EXPERIMENTAL HYBRIDS, SALINAS and IMPERIAL VALLEY, CA., 1998-99

(TESTS 199, 299, 2699, 3599, 5899, B399 and B899)

(PM-ERR)	ERR-DI	o-		3.7	•	0.3	0.7	20.8	8.4		9.0	1.0				10.5	22.6									
Test 3599	Powdery Mildew	o40 		5.1	5.0		6.3	•	5.3		6.1	4.9			7.0	7.1	(T)									
(NB-Line)	МО	% ∣		1.1		29.9	52.0	14.5	48.2	20.4	3.4	64.7	33.3				50.3(Iso)		•	33.0	8	32.0	41.2	34.4	41.8	43.8
Test 299 (N	Bolting	₩		32.5	0.0	27.7	20.0	43.5	41.7	_	3.5	_	45.7				41.5		56.3		33.5		23.0	33.2	40.0	33.4
(NB-Hyb)	MO	%		11.5	39.7	14.5	19.0	24.4	•	3.5	•	46.8	15.2		•	37.2	•	23.1	19.3	20.9	18.6	15.9	•	22.2	17.4	18.6
Test 199 (1	Bolting	o ∻ I	•	48.8	1.1	54.7	40.6	45.6	33.4	18.8	•	33.5	37.8		52.3	73.7	38.6	45.7	51.1	43.0	38.4	45.7	•	m.	46.3	41.9
1d)	RJAP	%	s (cont.	•	84.8	83.5	83.6	84.3	84.0	84.3	84.6	83.8	84.3	2 popus	84.0	84.5	84.7	84.1	83.8	84.5	84.4	85.0	83.8	85.2	83.3	84.0
t 2699 (yield)	Sucrose	∞ 1	Sf, Aa, Rz popns		16.25	16.39	17.14	16.63	17.08	16.56	•	16.68	16.65	from MM,S ^f ,Aa,R22	16.27	17.20	16.33	16.20	15.79	15.79	•	16.06	6.	•	6.2	15.86
Test	Sugar	1bs	from MM, VY, St		14082	14039	14236	14985	13970	12993	13639	15489	14137		13763	14558	14284	13128	13232	12854	12679	13724	13762	12596	13847	13306
	Variety			8929-41H50	8929-72H50	8929-102H50	8929-112H50	8929-114H50	8929-115H50	8929-133H50	8929-153H50	8929-154H50	8924H50	Lines & S1 pollinators	4035R	Rizor	х 869н50	R835H50	R836H50	Y873BH50	R879H50	X867H50	X872H50	X875H50	8926H50 (Sp)	8926H50 (Iso)

EVALUATION OF EXPERIMENTAL HYBRIDS, SALINAS and IMPERIAL VALLEY, CA., 1998-99

(TESTS 199, 299, 2699, 3599, 5899, B399 and B899)

(cont.)

	Test 2699	2699 (yield)	(g)	Test 199 (NB-Hyb)	NB-Hyb)	Test 299 (NB-Line)	NB-Line)	Test 3599 (PM-ERR)	(PM-ERR)
Variety	Sugar Yield	Sucrose	RJAP	Bolting	MQ	Bolting	DM	Powdery Mildew	ERR-DI
	lbs	æ1	de	%	% [%	₩	o%	o/e
Lines & S1 pollinators from MM, Sf, Aa, R22 popns	ators from M	M, S ^f , Aa, R22	sudod 3	(cont.)					
8927-29H50	14514	16.91	83.9	43.9	43.8	23.5	49.6	5.3	1.3
8927-30H50	13105	16.15	82.1	39.3	33.3	29.3	55.3	5.8	5.7
8927-33H50	13770	16.55	83.8	52.1	25.1	54.3	8.4	6.4	9.0
8927-37H50	14542	16.27	85.2	76.7	26.0	80.3	17.1	6.8	5.8
Mean	14041.7	16.46	84.2	38.5	23.7	33.1	31.8	6 .5	13.5
LSD (.05)	1321.4	0.62	1.6	15.5	18.9	17.3	19.8	9.0	6.8
C.V. (%)	9.6	3.84	2.0	25.0	49.5	32.6	38.8	6.1	40.6
F value	2.2**	6.17**	2.0**	6.5**	1.9**	10.1**	6.7**	25.5**	40.3**

EVALUATION OF EXPERIMENTAL HYBRIDS, SALINAS and IMPERIAL VALLEY, CA., 1998-99

(TESTS 199, 299, 2699, 3599, 5899, B399 and B899)

(cont.)

	Test	t 5899 (Rzm)	(m)	Test	B399 (IV-	(IV-Line)	Test	B899	(IV-Hyb)
	Sugar			Sugar			Sugar		
Variety	Yield	Sucrose	RJAP	Yield	Sucrose	Bolting	Yield	Sucrose	Bolting
	1bs	o%	% ∣	1bs	o%	%	lbs	o(0	o(c)
Experimental hybrids	٠.	- '73							
SS-432R	7983	17.73	84.1						
Rifle	8501	•	വ	11859	5.1	2.1	8270	13.35	
B4776R	10973	18.25	_	11034	15.49	0.0	9240	9. 9	0.0
8931H50	8323	16.45	82.7	11693	14.69	0.3	G	8	0.0
8925-19H50	10653	17.70	87.0	12336	14.05	0.0	9372	8	0.0
8913-70H50	9685	17.63	85.2	9488	13.44	3.4	8547	13.56	0.0
8911-4-10H50	10118	18.08	83.7	ω	ω.	8.0	15	2	0.0
8918-12H50	9651	17.83	88.1	10975	14.36	1.9	9286	•	0.0
8918-21H50	7672	17.00	•	10018		•	6609	•	0.0
Z825-6H50	8299	17.50	87.0	12388	14.99	1.9	8705	12.81	1.6
2825-9н50	10250	18.10	•	10829	15.60	0.3	7334	•	•
Z830-11H50	8023	16.50	85.1	13284	14.48	5.6	8321	12.10	0.0
R709-1H50	9039	17.48	84.2						
CR812H50	7163	17.40	84.0				7163	12.13	6.0
CR813H50	8613	16.65	5				8647	12.25	2.4
R709-9H50	8379	15.93	85.1						
S ₁ pollinators from N	from MM, VY, S ^f ,	Aa, Rz popns	នុក						
R878H50	2067	17.30	6.	11533	14.36	•	8478	•	1.9
8930-19H50	8103	17.35	85.9	12582	14.96	0.0	8569	13.26	0.0
8930-39H50	7617		4.	12030	14.85	•	8752	•	0.0
8930-102H50	7884	17.83	5.	10693	15.24	0.0	7575	•	0.0
R882H50	8790	16.65	85.7	10876	14.31	0.0	7334	12.58	0.0
R876-89-5H50			1		1	ı	8663	•	•

(TESTS 199, 299, 2699, 3599, 5899, B399 and B899)

(cont.)

	Test	5899	(Rzm)	Test	. B399 (IV-Line)	-Line)	Test	st B899 (IV-Hyb)	/-Hyb)
Variety	Sugar Yield	Sucrose	RJAP	Sugar Yield	Sucrose	Bolting	Sugar Yield	Sucrose	Bolting
	1bs	ov]	o,⊳	1bs	o⁄e	o/e	lbs	o/o	o∜∘
pollinators from	n MM, VY, S ^f ,	Aa, Rz popns	ns (cont.)						
8929-41H50	9948	17.13	85.0	12219	14.97	0.0	9257	3.5	6.0
8929-72H50	6260	16.88	•	11210	15.21	0.3	8841	13.29	
8929-102H50	8914	17.38	85.5	12258	4.6	1.2	8619	2.3	0.0
8929-112H50	10463	17.55		11503	15.19	•	9303	3.4	1.9
9-114H50	11174	17.88	86.6	11800	4.	0.3	8328	14.52	0.0
8929-115H50	8811	17.48	е Э	10919	15.47	1.5	0666	ά.	0.0
8929-133H50	9376	17.23	•	6710	5.6		7030	3.8	0.0
8929-153H50	7712	17.33	85.7	ω	14.79	0.0	8797	13.19	0.0
929-154H50	9209	17.58	83.6	10593	14.11	0.0	6518	1.6	0.0
8924H50	7335	16.75	86.7	10359	14.50	2.6			
Lines & S1 pollinators	from	MM, S ^f , Aa, R22	22 popus						
ir.	9144	17.55	œ				7593	12.78	0.0
Rizor	9416	18.05	85.4	12131	15.31	3.5	8820	•	0.0
X869H50	9054	16.90	85.1						
R835H50	8663	16.95	85.2				9969	12.94	0.0
R836H50	8834	16.83	85.4				6129	11.88	0.0
Y873BH50	1960	16.50	86.7				7595	12.75	0.0
R879H50	7003	15.13	86.5				7130	11.74	1.0
х867н 50	7366	16.70	85.0				8290	12.32	0.0
Y872H50	10374	16.45	84.7				7813	•	1.3
X875H50	8442	16.77	86.5				7458	2.	0.0

EVALUATION OF EXPERIMENTAL HYBRIDS, SALINAS and IMPERIAL VALLEY, CA., 1998-99

(TESTS 199, 299, 2699, 3599, 5899, B399 and B899)

(cont.)

'-Hyb)	Bolting	%				6.0	0.0	0.0	1.0	6.9	<	2.5	394.9	1.5*
Test B899 (IV-Hyb)	Sucrose	o/0				13.24	12.99	13.05	13.11	12.10	12 69	1.16	6.56	
Tes	Sugar	1bs				7927	7938	9141	7852	7751	7705.6	1574.8	14.7	3.0**
Line)	Bolting	~					2.3	8.0	5.6	16.8	6	3.5	182.8	**6.9
Test B399 (IV-Line)	Sucrose	o⁄o					15.28	15.35	15.26	14.46	14.90	0.87	5.92	1 2.95**
Test	Sugar	1bs	(cont.)				11030	11372	9280	11137	11178.1	1297.9	11.8	6.6 **
m)	RJAP	o⁄0		84.6	83.4		85.4	85.3	85.3	85.9	85.4	2.5	2.1	1.6*
Test 5899 (Rzm)	Sucrose	o,⇔	1, S ^f , Aa, R2	16.40	16.65		18.23	17.03	17.80	16.92	17.16	0.77	3.19	* 6.91**
Test	Sugar Yield	1bs	rs from M	8367	8705	ن)	8168	8156	9042	8628	8683.4	1938.5	16.0	2.7**
	Variety		Lines & S1 pollinators from MM, Sf, Aa, R22 popns	8926H50 (Sp)	8926H50 (Iso)	8926H50 (Sp or Iso?)	8927-29H50	8927-30H50	8927-33H50	8927-37H50	Mean	LSD (.05)	C.V. (%)	F value

(TESTS 399, 1399, 4399 and B1299)

	T.	Test 399 ((NB)		Test	Test 1399 (VY)	X)		Ě	Test 4399	(Rzm)		Test B1	Test B1299(IV)
	1	Stand		Sugar		-	Beets/	ΜĀ	Sugar			Root	Appear	%Liv
Variety	%Bolt	Count	ΩM	Yield	Sucrose	RJAP	1001	Score	Yield	Sucrose	RJAP	Rot	Score	Plants
	8/26	Mean	o∤≎ [1bs	o⁄∘	ονΙ	No.	Mean	1bs	æ 	o ∤ 0	%	8/2	1/8
Checks US H11	19.4	17.0	14.1						4718		83.7	25.4	4.5	21.9
) }								5579	ω.	87.1	11.1		
									4073	ლ. ი დ. ი	9 9 9 0 1	16.3		(
									5179	χ. Σ	83.5	0.0	4.0	13.3
US H11 USH11													4.5 0.0	5.3 0.0
SS-NB3		16.0	27.2											
B4776R		16.3	31.9						8636	17.27	87.5	7.5	3.0	60.1
97-C37		17.0												
97-SP22-0		17.3	40.3	4491	14.70	83.2	164	6.9						
97-US22/3	•	15.0												
R853	15.9	16.3	4						2600	15.57	84.6	17.3		
X873	•	16.0	33.9						8233	16.27	83.3	21.0		
X867	45.4	17.0	47.0	8884	16.63	84.7	145	5.0	7301	16.33	94.6	17.5		
X867													1.0	73.1
X872	25.4	15.7	42.8						8928	16.27	84.0	0.0		
¥872													1.5	67.6
R836									5972	14.63		11.1		
R879									5526	15.20	85.1	30.4		
R840									7823	16.20		3.6	L	
Kitle													n c	•
55-776K) -	•
													1.0	
													3.5	
													2.5	
													3.0	30.2
Y875 (Iso)													о· В	

(TESTS 399, 1399, 4399 and B1299)

	Ē	Test 399 ((NB)		E. 50	1399 (V	(XA)		Ĭ	Test 4399	(Rzm)		Test B1	B1299(IV)
				Sugar		-	Beets/	ΔĀ	Sugar			Root	Appear	%Liv
Variety	%Bolt	Count	DM	Yield	Sucrose	RJAP	1001	Score	Yield	Sucrose	RJAP	Rot	Score	Plants
	8/26	Mean	%	lbs	%	% ∣	No.	Mean	lbs	₩	%	%	8/2	1/8
R846-# = RZM	R746PX	= c37*3 x	r R22											
R846 - 1	9.6	16.3	33.1	6369	13.90	82.2	155	6.0	6858	14.20	82.2	7.5	•	$^{\circ}$
- 2	12.4	16.3	41.2	6182	Ŋ	т М	3	•					3.5	38.1
د ا	15.5	15.7	53.8						7918	15.30	84.4	5.3	•	0
- 4	35.1	15.3	38.7										•	13.6
ı	17.1	14.0	67.5											
9 1									7186	5.3	9	•		•
- 7									6677	14.80	81.9	1.9	3.5	19.6
& 1									7432	5.1	4	•		
R853-# = RZM	R753PX	= C37*4 x	r R22											
1	15.4	6.		6490	5	9	155	•	6693	4.	86.4	н.	•	
- 2	16.7	16.3	64.6	6754	15.43	86.1	124	4.9	6422	15.03	85.1	13.0	3.5	31.3
ю 1	12.5	9		7286	Ω	4.	118	5.3	6637	Ŋ.	84.8	•	•	•
- 4	23.9		70.5	6667	9	ö.	$^{\circ}$	•	6408	Ω.	کا	•	•	4.
I U	17.1	16.0	64.7	8796	Ω	9	2	•	5760	æ.	'n	•	•	ė.
9	20.8	15.3	70.8	8345	15.60	83.0	148	5.4	9029	5.1	ω.	2.4	•	
- 7	0.0		•	7057	5.4	5	5	5.4	7899	5.0	5	•	•	Ŋ.
& 1	0.0	15.7	54.6						5361	15.30	82.9	37.5	4.5	14.3
ი 1	•	•	28.2						7458	5.3	4.	•	•	。
-10		•	•						7023	4.5	9	•	•	5.
-11	5.1	14.0	27.2						7104	5.3	4	•	•	
-12	6.8	14.7	49.7						6498	5.3	ω	8	•	•
-13	40.0	14.7	51.3						6543	14.83	84.3	20.7	3.5	41.3
-14	27.6	14.3	32.9						9057	5.6	4.	0	•	•
-15	7.2	14.3											•	•

(TESTS 399, 1399, 4399 and B1299)

	Ē	Test 399 ((RR)		E A	Test 1399 (VY)	2		E	Test 4399 (Rzm)	(Rzm)		Tost 131	Test R1299(TV)
				Sugar			Beets/	ΛX	Sugar		,	Root	Appear	%Liv
Variety	%Bolt	Count	DM	Yield	Sucrose	RJAP	1001	Score	Yield	Sucrose	RJAP	Rot		Plants
	8/26	Mean	æ	lbs	o, € [o(0	No.	Mean	lbs	∞ 1	%	%	8/1	8/1
R853-# = RZM	R753PX	= C37*4 x	R22	(cont.)										
R853-16	5.	S	۲.										4.0	9
-17	13.7	17.0	27.5										•	80
-18									7002	4.	4	•	•	ω.
-19									7294	15.57	86.8	2.1	3.5	31.4
-20													•	<u>و</u>
-21													3.5	5
22- 17:													4.5	14.3
3														
X873-# = RZN	RZM Y773 PX	ıı	$7 \times Y71$											
Y873 - 1	7	9	<u>ი</u>	6100	15.40	4.	136	•	5761	15.5	9	5	•	•
- 2	64.6	9	7.	8828	15.97	2	109	•	6623	15.9	Η.	•	•	4.
m I	37.0	14.3	31.3	5600	14.67	82.4	124	5.8	6547	15.57	82.5	12.4	3.0	43.3
- 4	55.1	5	δ.	7230	15.17	4.	127	•	7824	15.6	5	•	•	9
ı S	33.1	15.3	54.3	5380	15.30	ω.	142		4660	16.1	و	•	•	ω.
9		5	ω.	7225	14.97	ω.	133	•	7625	5.3	4.	14.5	4.0	ω.
- 7	•	16.3	38.8	6430	15.83	82.9	152	5.5	6320	15.93	84.6	1.8	4.0	34.2
8 1	41.4	5	7.	6965	15.20	4.	142	•	7746	5.7	7	2.2	4.5	•
ი I	24.6	9	•	7565	16.33	4.	145	•	7310	5.4	e.	•	•	
-10	9	4.	5	6757	15.47		139	•	7197	5.7	4.			ω.
-11	4.	5.	ъ.	6367	15.10		133	5.8	7583	5.5	∺.			
-12	59.3	15.3	24.2	ω	4			6.1	5008	14.17	85.8	13.1	4.5	7.1
-13	<u>ი</u>	5	ω.						7896	5.7	w.			ი
-14	m.	5.	ω.						6484	5.4	æ.		•	
-15	7.	2	Ŋ.						7747	6.3	4.			7.

(TESTS 399, 1399, 4399 and B1299)

Stand Stand Name Tase 1 1399 (WY) Store Stand	84.8 8.5 3.5 82.9 4.2 1.5 85.0 8.6 2.0
Name	5.0 8.
Name	4.27
Comparable Court	
(NB) Sugar Beets/ VY Sugar Sugar Sugar Beets/ VY Sugar Sugar Sugar Sucrose RJAP 100' Score Yield Sucrose RJAP 100' Score Yield Sucrose RJAP 100' Score Sield Sucrose RJAP 100' Score Sield Sucrose RJAP 100' Score Sield Sucrose Sield Sie	15.83 15.73 14.40
NB Sugar Beets	9436 8090 5499
NB Sugar Beets	7. 7. 4. w. w. w.
NB Sugar Sugar Sugar Sugar Sugar PM Yield Sucrose RJAP Sugar S	148 136 155
x (O.P. x R22) 21.4 8938 16.27 21.4 8938 16.27 36.9 7870 16.80 59.1 8835 16.23 34.3 10580 17.83 57.4 9579 16.93 37.9 10371 17.00 31.8 x R22	85.5 84.0 81.7
(NB) Sugar PM Yield \$ 1bs 100.0 58.7 100.0 58.7 21.4 8938 36.9 7836 45.5 7870 59.1 8835 34.3 10580 30.0 8620 57.4 9579 33.8 8625 37.9 10371 31.8 * R22	16.43 16.00 17.57
(NB) DM 8 8 137 x Y71) 35.7 100.0 58.7 58.7 21.4 36.9 45.5 59.1 34.3 30.0 57.4 33.8 31.8 x R22 x R22 x R22 33.5	9195 9777 7252
<u>∀</u> <u>'</u>	50.0 57.9 51.7
	16.0 16.0 14.3
	56.3 14.8 35.8
RZM RZM	1 1 1 ധെ 4 സ

(TESTS 399, 1399, 4399 and B1299)

	Τ¢	Test 399	(NB)	0	Test	Test 1399 (V	(VY)		Ţ	Test 4399 (Rzm)	(Rzm)		Test B	Test B1299(IV)
Variety	*Bolt	Stand	ΝC	Sugar	Sucrose	R.TAP	Beets/	VY	Sugar	Sucrose	R.TAP	Root	Appear	%Liv
	8/26	Mean	 	1bs		o(0	No.	Mean	1bs	% %	o(0	o o o l	7/8	7/8
Y871-# = RZM	4 Y771 PX	= 0.P.	x R22 (c	(cont.)										
1		٦.	55.3	9432	6.4	5	136	•	7	5.9	5	•	•	ω.
- 7		4	36.6	7879	15.77	83.0	133	5.3	ന	4.7	4		•	ω.
& I	57.8	15.0	15.6	5679	6.1	w.	115	•	5905	14.33	79.5	37.5	•	7.
ი I	•	8	26.2						0	6.1	ж Э	•	•	5
-10	•	14.0	4.						œ	4.2	2	•	3.5	30.8
-11													•	0
X872-# = RZM	4 Y772 PX	= R80,R7	76 × (C37	7 x R22)										
X872 - 1	70.7	15.0	4.		16.	ъ.	Н		7227	15.77	4.		•	
- 2	4.	•	9	7118	15.	4	Н		œ	5.	H.		•	•
m I	26.1	15.3	5	10456	15.	5	2	•	8565	4	5	7.8		
- 4	6.3	15.7	36.5	6441	16.87	83.0	115	6.3	8601	16.90	82.5	0.0	1.5	64.7
េល	32.2	15.7	9.	8589	9	7.	0		7268	6.4	8	20.7	•	•
9	0.0	13.3	55.3	6291	رح	∺.	М	•	40	6.0	2	•		₽.
- 7	18.7	16.0	4.	7799	15.57	82.2	118	6.2	59	6.0	m.	0	•	근.
6 0 1	43.3	14.7	35.7	8613	Б.	ъ.	Н	•	8266	16.23	86.8	17.3	•	6
ი 1		•	2						85	6.3	8		1.0	74.8
-10													•	4
#	R776-89-5NB	X RZM	7934-# (C	(C913-70aa	x R636)									
													•	0
- 2													•	•
m I													•	9
- 4													•	5.
I I													1.5	74.4
o I														7.

(TESTS 399, 1399, 4399 and B1299)

Variety %Bolt 8926-# = RZM 7926⊗ 8926 - 1 8926 - 1 - 2 - 3 - 4 - 5 - 6 - 6 - 7	Stand %Bolt Count DM 8/26 Mean % 926⊗ = MM,S [£] ,Aa,R22	DM 	Sugar Yield	ose	:	Beets/	λλ	Sugar	1	(17.71)		Appear %Liv	(17)667
Wariety 8 8926-# = RZM 79 8926 - 1 - 2 - 3 - 4 - 5 - 5 - 6			- 1	Sucrose	5							15) 44.	%Liv
8926-# = RZM 79 8926 - 1 - 2 - 3 - 4 - 5 - 5 - 6 - 7			lbs		RUAE	100'	Score	Yield	Sucrose	RJAP	Rot		Plants
8926-# = RZM 8926 - 1 - 2 - 3 - 4 - 5 - 5 - 6 - 7				o 0	o/0	<u>ا</u> ۋ	Mean	lbs	op]	op	æi	8/1	1/8
8			(4)										
												2.5	72.9
												3.0	43.3
1 1 1 1 1												4.0	18.4
1 1 1 1												3.0	0.09
1 1 1												5.0	0.0
1 1												5.0	
												4.0	•
ı												0.0	0.0
- 1-1												0.0	•
01												4.5	
-11												4.0	14 6
-12													
-13												Ն Մ	24.5
-14) L	7 .
-15												, r.	
-16												, c	•
												, ,	•
= R778%	x RZM P707B	((Y71 x P	603) (~CP	P603) (~CP01)) (gh 10)	ଚା								
Z = Z = Z												•	74.1
7 * I												2.5	61.6
ភ (•	55.6
000 I												•	9.99
P808B-# = R778% x	x RZM P708B	$((Y71 \times P)$	P604) (~CP02))	02)) (gh 10)	(0								
P808B- 2												2.5	63.1
τ) «												4.0	5.4
व ं ।												3.5	30.5
` 1												1.5	56.9

(TESTS 399, 1399, 4399 and B1299)

	Te	st 399 (1	(B)		Test :	Test 1399 (VY)	Y)		Te	Test 4399 (Rzm)	(Rzm)		Test B1299(IV)	299 (IV)
		Stand		Sugar			Beets/	Λλ	Sugar			Root	Appear	%Liv
Variety	%Bolt	Count	DM	Yield	Sucrose	RJAP	1001	Score	Yield	Yield Sucrose RJAP	RJAP		Score Plants	Plants
	8/26 Mean	Mean	oko	1bs	æ	o(0	No.	Mean	1bs	ok∘	ok∘	æ1	8/1	8/1
Mean	39.5	15.2 4	40.4	7692	7692.9 15.91	83.9	135.9	135.9 5.5	7319.	7319.3 15.45	84.4	84.4 10.9 3.2	3.2	37.1
LSD (.05)	22.0	1.9	34.2	1626	.6 0.73	3.0	33.1	9.0	199	1991.8 1.01	3.7	3.7 20.5 1.4	1.4	32.8
C.V. (%)	34.6	7.9	52.5	13	.0 2.83	2.2	15.0	15.0 6.8		16.9 4.06 2.7 116.6 22.8	2.7	116.6	22.8	44.7
F value	11.1**	8.8**	8.8** 1.9**	9	6.2**9.23**	2.5**		1S 5.8**		1**5.43**	1.9*	* 1.5*	4.7**	3.5**

canopy level. Due to root rot, high levels of bolting, and trimming, the second and third counts were more difficult to make. In general, it appears that counts for bolting made 8/26/99 are best indication of relative bolting tendency. Tests 199 thru 999 were infected with Sclerotium rolfsii, southern root rot. After each counting for bolting, seed stalks were trimmed to Level of bolting in 1999 tests was very high due to a long vernalization period in the winter and spring. See Tests 1399 and 4399 for performance under virus yellows and rhizomania.

Emergence scored on a scale of 0 to 5 where 0 = no emerged plants. Downy mildew infected plants counted 5/26/99. Downy mildew infection became moderately severe and probably affected bolting tendency and late summer survival. See tests 399 and 4399for companion tests under bolting and rhizomania conditions. Inoculated with VY (BYV-BWYV-BChV). TEST 1399 NOTES:

Rotted roots weighed but not included in sugar sample. Weights adjusted for missing feet, but not for TEST 4399 NOTES: Harvested by machine with 10% tare used. Root rot due to Sclerotium rolfsii. Rotted roots counted Also see results from bolting, virus yellows, and Brawley trials. before harvest. root rot.

TEST B1299 NOTES:

3 = intermediate and variable; 4 = fair; and 5 = poor to mostly dead plants. Appearance scored relative to the overall Appearance scored on a scale of 1 to 5 where: 1 = very good canopy; 2 = good canopy and appearance often segregating; assumption was that plant health and appearance was mostly being influenced by reaction to rhizomania and rhizomania under high temperature conditions. However, other factors such as plant vigor, cyst nematode infection, root rots, test at time and based upon canopy size, uniformity, color, vigor, number of dead leaves, and dead plants. The etc. could have influenced appearance.

Coefficients of correlation for % Living plants vs. Appearance scores for 5/12,6/11, & 7/8 and Stand Counts (October Stand counts made post thinning in October. 1998) are r = -.60**, -.72**, -.87**, and 0.01, respectively. plants counted 08 July 1999.

EVALUATION OF PAIRCROSSES (FULL SIBS) WITH C31Rz GERMPLASM, SALINAS, CA., 1999

(TESTS 499, 1499 and 4499)

	Te	Test 499	(NB)	11.	Test	1499 (VY)	Y)			Test 4499	(Rzm)	
		Stand		Sugar			Beets/	Λλ	Sugar			Root
Variety	%Bolt	Count	DM	Yield	Sucrose	RJAP	1001	Score	Yield	Sucrose	RJAP	Rot
	8/26	Mean	% 	lbs	∞	% [No.	Mean	lbs	₩	% 1	₩
Checks												
Y869 (Iso)	58.1	16.7		8799	6.0	m.	136	3.8	7054	ဖ		19.7
R876-89-5NB	21.4	17.3	5.8	9425	16.47	85.2	155	4.4	6577	16.80	ж Э	1.9
97-C37	3.9	16.7		7225		Э.	161	4.4				
97-SP22-0	82.8	15.3	ъ.	3149	3.1	•	133	6.4				
US H11									5223	3.4	4	•
B4776R									10547	17.10	86.7	5.3
Y868-# = RZM Y768	PX											
Y868 - 1	0.0	16.0	Н	8282	7.3	5	2	•	6934	7.	86.2	11.7
- 2	20.7	15.7	39.0	11354	17.00	85.5	145	3.8	7513	16.07	84.6	5.1
м 1	70.0	•	4	9423	6.8	æ.	4	•	7369	ø.		4.5
- 4	31.6	•	0	7831	6.9	5.	m	•	8258	•		4
ا ى	6.7	15.3	9	7220	4.6	•	2	•	6827	15.07	84.9	21.5
9	39.3	15.3	15.1	9753	16.83	82.5	4		7997	16.87	96.6	17.1
- 7	6	16.7	43.9	9916	16.47	86.7	139	4.3	7032	5	85.4	5.6
8 1	19.1	•	57.9	7429	5.0	Ŋ	ω	•	7120		9	9.4
6 1	74.1	14.3	7						7238	•	83.9	2.4
-10	9.3	14.0							6537	16.00	e.	2.4
-11	55.9	13.3	57.3						7540	ري		55.2
-12	7.5	13.3	54.8						6347	•		رى
-13	23.1		32.1						7788	9	88.2	12.5
-14	9.3	15.0	77.2						6047	ė.	•	8.3

(TESTS 499, 1499 and 4499)

(cont.)

	Root	Rot	%		4.4	4.9	16.6	•	•	7.7	14.1	2.0	15.0	4.3	2.4	•	6.3	32.9	11.9	4.1		•	12.0	•	o,	31.3	•	•
(Rzm)		RJAP	% l		m.	Э.	85.0	9	7.		84.1	84.3	85.2	84.8		4.	83.7	86.5		85.7	9	•	83.7	85.8	4.	85.1	•	85.6
Test 4499		Sucrose	% ∣		6.5	6.8	16.43	. 2	16.37	9	17.67	15.93	16.50		16.50	15.47	.5	15.20	ω.	16.00	ι.	6.9	16.30	5.7	17.27	5	რ.	5.3
	Sugar	Yield	1bs		9153	7042	8540	7905	7424	2	8550	47	7569	7818	7495	7090	8468	9	7297	_	0	m	2066	4	9	8196	4	8
	ΛX	Score	Mean		3.7	4.4	5.1	3.8	3.6	•	4.1	4.0	•	3.7	4.8	4.3	4.2	4.3	4.4	4.1	3.3	•	3.8	•	4.0	4.9	•	4.0
χ)	Beets/	1001	9		2	B	139	ന	4	124	148	145	3	148	158	136	139	124	136	112	124	121	115	118	r	109	E	3
1499 (VY)		RJAP	%		83.8	4.		84.1	84.3	2	82.4	ъ.	85.7	83.8		83.3	83.1	ω.	83.9	•	7.	0	84.9	ω.	83.3	9	83.7	
Test		Sucrose	% I		ė.	•	15.80	5	6.1	5.7	16.50	5.2	16.07	6.5	15.93	5.6	17.00	15.87	•	9	6.1	6.8	16.17	5.0	6.5	15.40	6.9	5.8
	Sugar	Yield	1bs		9892	8500	9653	92	11830	8421	8846	10088	10894	9931	10954	9177	10982	8116	Ŋ	9569	9357	10415	8968		8648	9336	10863	8883
(NB)		ΜΩ	∞		59.2	32.4	43.1	39.4	42.6	•	5.5	37.6	29.7	85.9	30.8	29.1	61.1	Э.	48.5		0	58.8	63.7	7	69.7	21.0	16.7	55.8
Test 499 (Stand	Count	Mean		16.7	ů.	16.3	7 .	15.7	9	17.0	7 .	•	16.7	18.0	7		15.3	16.0	5	14.7	•	16.7	ت	•	13.7	5	4.
Te		%Bolt	8/26	Xd 69	70.2	16.4	53.2	•	74.7	•	32.5	•	33.2	10.2	3.6	•	34.5	63.6	7.1	48.0	72.7	49.1	70.5	50.6		38.3		•
		Variety		Y869-# = RZM Y769		1 2	د ا	- 4	ı,	9 -	- 7	& 1	თ I	-10	-11	-12	-13	-14	-15	-16	-17	-18	-19	-20	-21	-22	-23	-24

EVALUATION OF PAIRCROSSES (FULL SIBS) WITH C31Rz GERMPLASM, SALINAS, CA., 1999

(TESTS 499, 1499 and 4499)

(cont.)

	Root	Rot	o%		4.4	2.0	7.0	23.1	•	3.2		0.0	0.0	5.1	•	0.0	•	•	•	•	2.2	•	•	0.0	•
(Rzm)		RJAP	%		υ	•	86.2	•	2	82.7	ω.	4.	79.1	ω.	84.4	ω.	84.0	4.	w.	6	81.8	ω.	Ή.	85.0	5.
Test 4499		Sucrose	% 		٥.	9	6.9	•	16.33	٠4	0.	7.3	15.90	ĸ.	16.97	•	7.9	9.	6.8	6	16.37	8.	6.9	17.33	7.1
	Sugar	Yield	1bs		8337	7672	7913	8071	5914	22	54	96	2608	67	80	7212	4	5480	7113	5256	6097	6857	79	7511	85
	Ž,	Score	Mean		4.6	9.8			4.3	4.6	•	4.1	4.3	4.3	4.3	4.7	3.9	•		4.2	4.4	4.2	•	4.2	•
(X.	Beets/	1001	No.		142	S			155	161	Ŋ	S	118	m	152	148	152	155	Ŋ	4	155	Ŋ	വ	142	2
1499 (VY)		RJAP	∞1		4.	82.5			84.0	•	•	2	81.3	•	•	82.6	•	Ή.	ω.	•	80.9	•	•	$^{\circ}$	•
Test		Sucrose	%		16.47				15.53	6.4	5.5		16.27			16.60		6.3	5.6	6.2	15.63	6.1	5.8	16.17	6.2
	Sugar	Yield	1bs		11479	10764			7835	L)	16	82	7899	56	7319	7866	8418	7062	8767	7825	7317	7825	10700	7575	8188
(NB)		MO	% 		2	Ŋ.	33.5	9	4.5	•	5.9	4.	14.3	Ŋ.	20.5	6.5	0.0		•	•	18.6		0.0	0.0	0.6
Test 499	Stand	Count	Mean	·	15.3	•	13.7	2	15.3	9	16.3	9	16.0		16.3	15.3	15.7	4.		υ.	15.7	9		14.7	
Te		%Bolt	8/26	PX (cont.)	7		62.8	•	27.0	10.1	24.1	29.5	0	43.3	61.5	37.2	28.6	•	•	ω.	35.5	•	Η.	31.6	7
		Variety		Y869-# = RZM Y769	\sim	-26	-27	-28	R886-89-5NB- 1	1 2	en I	1 - 4	I G	9 1	- 7	80 I	6 I	-10	-11	-12	-13	-14	-15	-16	-17

(TESTS 499, 1499 and 4499)

(cont.)

	Tes	Test 499 (1	(NB)		Test	Test 1499 (VY)	Y)		1	Test 4499 (Rzm)	(Rzm)	
		Stand		Sugar			Beets/	ζ	Sugar			Root
Variety	%Bolt	Count	DM	Yield	Sucrose	RJAP	1001	Score	Yield	Sucrose	RJAP	Rot
	8/26	Mean	%	1bs	æ	%	No.	Mean	1bs	&	%	%
R876-# = RZM R776	PX											
	32.3	15.7	15.0	9404	5.8	m.	136	4.8	7750	.2	4.	•
- 2	33.1	ъ.	30.0	7022	14.80	84.3	130	5.2	7136	15.57	85.4	2.1
ო 1	0.0	16.0	57.7	7803	5.2	4.	127	5.0	7475	9.	9	•
- 4	13.1	9	30.5	8028	4.6	e.	142	•	7328	5.6	4.	•
l R	2.1	16.0	41.7	4	5.1	4.	124	4.4	51	ω.	5	4.8
9 1	29.9	9	16.7	σ	6.3	4.	145	4.3	10	6.2	4	
- 7	4.2	16.0	43.8		15.40	82.7	145	5.0	7818	15.33	83.3	1.9
80 I	10.3	9	14.2	24	5.1	ъ.	133	•	96	5.2	4.	
c I	c	u	_	8103	L L	~	צו	r m	6492	~	_	
	•		1 () () () L) (•	י נ			•
-10	4.	ė	2.69	T0623	ე ა	ດ .	η,	٠	9	٥. ر	84.4	٠
-11	22.5	16.3	53.1	7286	15.10	83.6	124	4.7	7266	15.47	84.2	0.0
-12	45.8	رى	17.2	8739	6.0	ъ.	2	4.8	52	6.2	86.2	•
R881-43-# = RZM R781	781-43 PX											
R881-43 - 1	5.9	9	Н	44	0.	ک	4	5.8	51	5.3	5.	•
- 2	26.8	9	σ	9139	ω.	ė.	r	•	20	6.5	5.	•
ო 1	0.0	15.3	ß	7481	ω.	4.	$^{\circ}$	•	91	5.8	7.	
- 4	•	•	34.1	7974	15.67	82.7	145	5.2	7535	16.07	84.0	2.0
ı ا	47.3	14.3	ဖ	10652		ъ.	Н	•	85	6.0	7	•
ı	1 80	7	92.4	8545	9	4	Ľ	7 7	76	15,67	86.4	
			I C	7317	ו ר		_		ָ '	· u	. u	
	•	• [1 t) L	, (۳ <	•	1 0		o (
	62.7	17.0	90.8 10.8	1261	15.60	92.7	140 100	0 L	1087	15.90		0 0
ე 	٠	7	m.	8.745	ა 1	· ·	າ)	•	7	٠	83.4	
-10	14.7	16.7	27.3	8622	6.0	m	ന	•	92	ė.	4	

EVALUATION OF PAIRCROSSES (FULL SIBS) WITH C31Rz GERMPLASM, SALINAS, CA., 1999

(TESTS 499, 1499 and 4499)

(cont.)

	Te	Test 499 ((NB)	11	Test	1499 (VY)	X)			Test 4499	(Rzm)	
		Stand		Sugar	i		Beets/	ĀΛ	Sugar			Root
Variety	%Bolt	Count	DM	Yield	Sucrose	RJAP	1001	Score	Yield	Sucrose	RJAP	Rot
	8/26	Mean	% 1	lbs	.e%1	% 1	No.	Mean	1bs	%	o(0	ol i
R881-# = RZM R781	PX											
	•	5	9	10183	4.	ω	148	4.0	7095	4.	5.	•
- 2	71.8	14.3	46.5	10402	15.47	83.4	148	4.5	7966	15.60	84.9	2.1
г Э	•	ė.	Н	10827	4.		158	•	9790	5	5	•
- 4	61.4	7 .	_	11115	9	5	4	4.7	7604	9	•	•
. 5	48.8	15.0	53.0	8611	15.27	84.6	148	5.3	04	5.5	4.	4.2
9 -	•	9	34.9	10186	ъ.	84.5	142	4.1	50	•	85.9	•
- 7	65.5	15.0	47.9	8439	9.	82.7	158	•	9042	5.3	7.	2.1
80 1	64.0	9	•	8604	2	83.4	161	5.3	30	4.	83.7	16.4
6	8	9	0	9002	5.6	4.	വ	5.1	8918	6.1	7.	7.4
-10	65.5	•	\vdash	11175	15.70	83.5	4	4.4	9454	15.73	84.9	9.8
-11	42.9	16.3	63.2	9704	4.8	•	136	•	9117	ъ.	4.	•
12	41.8	9	2	8884	14.93	84.4	9	4.8	9184	14.33	85.8	9.4
-13	56.5	5	0	9552	5.1	4.	2	4.1	9675	5.2	4.	3.7
-14	67.4	9	42.5	9427	6.1	5	m	5.0	8	5.6	85.5	10.6
-15	29.6	15.7	47.9	8664	15.23	84.7	139	4.5	8439	15.53	•	•
-16	0	•	19.6	9018	5.2	4.	N	4.6	31	5.9	86.4	7.5
-17	61.2	9	8	7324	\mathbf{c}	81.8	m	4.8	9400	16.03	5	9.6
-18	ά.	15.7	74.3	10383	ന	5	$^{\circ}$	4.8	9829	15.00	85.7	11.8
-19	53.6	ر. د	40.7	7	15.00	Ŋ	2	•	N	5	5	
-20	38.9	9	σ	11893	S	82.8	124	•	8239	15.17	84.0	8.9
-21	35.2	13.7	71.3	7189	14.87	83.5	0	4.8	6889	15.03	85.3	9.8

(TESTS 499, 1499 and 4499)

(cont.)

	Root	Rot	₩			23.5	29.8	13.7	4.	ω.												8.6	16.6	120.1	* 2.7**
(Rzm)		RJAP	o⁄e		86.0	84.5		83.4	83.3	2												84.9	2.9	2.2 1	2.2**
Test 4499		Sucrose	%		16.00	17.10	16.97	15.83	5.2	5.5												Н	3 0.91	5 3.51	4** 6.02**
	Sugar	Yield	1bs		6246	9083	8787	9198	5154	6450												610.	١٠.	ъ.	2.,
	۸X	Score	Mean		4.0	4.8	4.6	4.4	4.7	5.1			5.2	4.5	4.4	4.0		4.2	4.9	4.6	4.7	4.5	9.0	7.9	* 6.5**
C	Beets/	1001	 		139	136	145	142	145	142			121	142	130	139	,	145	161	148	155	ത	28.0	12.5	٠. *
1499 (VY)		RJAP	%		85.2	82.7	83.0	84.0	82.7	84.7			•	80.9	•	•	,	83.5	81.7	83.1	86.7	83.9	2.9	2.1	1.8**
Test		Sucrose	. %		16.07	16.03	16.80	15.97	6.3	6.0			Ŋ.	15.87	ø.	16.20		رى	15.40	Ŋ.	ø.	15.80		3.17	*
	Sugar	Yield	1bs		7686	9047	10426	10974	8465	8257			7757	8881	8746	10211		9409	9346	9962	9137	m	_	13.6	w.
(NB)		DM	ok∘		61.7	33.3		57.6	37.9	39.6							ts					38.4	30.6	49.6	
499	Stand	Count	Mean		15.3	16.0	16.7	15.7	16.7	17.0							ing Tests					15.8	1.9	7.6	2.1**
Test	i	%Bolt	8/26	0 PX	4.4	37.5	52.1	64.0	0.0	29.6		g Test 599					ded in Bolt					38.1	22.5	36.8	7.7**
		Variety		R870-# = RZM R770	R870 - 1	1 2	_ا ع	- 4	ا د	9 -	=======================================	R8/0-# in Bolting	R870 - 7	80 1	ი 1	-10	R870-# not included in Bolting	R870-11	-12	-13	-14	Mean	LSD (.05)	C.V. (%)	F value

EVALUATION OF PAIRCROSSES (FULL SIBS) OF C78 & C80, SALINAS, CA., 1999

(TESTS 599, 1599, 4599)

9 (Rzm)	RZM Root	AP Resist Rot	&	1 90.3 6	93.4 10.							.0 85.2 8.		.9 96.1 3.	.3 83.9 15.	.0 85.9 7.	.4 86.7 4.0		4 87.5 11.	1.7 17.		4 84.6 0.	8 26	3 96.1 16.		.3 32.5 0.0
Test 4599		Sucrose RJAP	o, € [53 84	6.70 83							6.63 85	17.03 83	6.97 85	6.77 86	6.80 83	77 84		6.03 83	82	7.63 84	6.70 83	97 84	7.13 83	((15.63 82.
	Sugar		1bs	8131	7922							8807	7825	8147	8782	7774	8126	,	8922	8031	8817	7742	8527	8215	(3093
	ΛX	Score	Mean	5.0			0.9			4.6		•	4.8	•			4.8		٠	4.9	•	•	4.8	•		٥.
VY)	Beets/	1001	No.	167	139	152	167	148	167	152		4	148	2	വ	4	136		m	139		4	145	2	(LJy
1599 (VY)		RJAP	% ∣	84.9	4.	Η.	82.3	4.	ά.	ά.		9	82.8	5.	ى	Η.	84.7		4	\sim	82.9	۲,	84.3	ω.		82.0
Test		Sucrose	o4∘ I	16.43	6.2	4.9	15.17	6.1	6.4	5.7		6.2	16.73	6.7	6.5	6.4	16.57		5. 8	16.00	7.0	5.9	16.30	6.5	(16.00
. 13-	Sugar	Yield	lbs	8593	7633	4605	3087	9251	7795	8573		9206	9108	7668	9476	6502	8347	,	8439	7817	7255	7662	8858	8849		5004
(NB)		DM	æI	33.4	49.4							37.7	58.5	50.8	44.6		77.1		77.1	45.1	6.99	84.1	55.7	68.89		7.7/
Test 599 (Stand	Count	Mean	17.3								16.0	16.0		17.3	9			16.0	17.0	16.3	16.0	9	4	,	D.0T
Тe		%Bolt	8/26	32.1	30.4						R778 PX	95.4	58.5	4.2	9.6	18.0	62.5		•	2.0	•	12.4	•	30.9	R780 PX	0.8
		Variety	مامولان	R878%	R880	97-US75	97-SP22-0	R881	R876-89-5NB	Y869 (Iso)	-# = RZM	ı	- 2	۳ ۱	- 4	1	9	ı	- 7	8 0 І	ი I	-10	-11	-12	щ.	T = noov

EVALUATION OF PAIRCROSSES (FULL SIBS) OF C78 & C80, SALINAS, CA., 1999

(TESTS 599, 1599, 4599)

İ	Root	Rot	ok∘ I		•	•	0.0	•	7.4		7.8	•	Ή.	27.8	ö	•	о О	•	9.3		•	16.9		•	•	15.7	•	
(Rzm)	RZM	Resist	oko		•	6	93.9	ω.		4.	89.2	2	ω.	78.5	Η.	2	•	7.	97.9	o.	ω.	87.1	ı	٠.	9	78.3	7	ω.
4599 (1		RJAP	o∤e		4.	2	83.0	ë.		8	83.8	5.	ω.	83.9	2	•	Ή.	4.	83.7	т М	82.1	m		ď	4.	84.2	4.	4.
Test		Sucrose	de		9.9	6.0	17.30	6.7	5.9	6.4	17.30	5.9	5.7	16.33	7.3	6.3	4.8	ω.	17.10	16.83	16.53	6.5	1	9	6.8	16.50	7.5	6.5
İ	Sugar	Yield	1bs		8320	5601	8740	6202	7708	8069	12	\vdash	8054	7038	7210	6572	5613	8519	7228	7645	8380	78	(ρ	8611	7516	8592	78
	ΛX	Score	Mean		•	•	5.6	•	4.3	•	5.2	•	•	•	5.2	•	4.8	•	4.3	•	5.0	•	(5.2	5.1	5.3	5.3	5.2
(VY)	Beets/	100	No.				142		9	2	152	9	c	142	2	155	148	139	133	139	124	124	,	136	142	142	139	130
1599 (\		RJAP	o(•		5.	Э.	84.2	2.	0	ω.	83.6	ω.	ω.	82.5	Э.	8	4	ω.	81.8	4.	83.5	2		'n	ო	85.8	ω.	4.
Test		Sucrose	%		9	15.77	ဖ	16.33	5.9	4.	9	5.9	5.7	16.67	6.8	15.97	5	•	16.20	5	15.37	5	(ų.	ė.	16.97	ė.	9
	Sugar	Yield	1bs		81	71	51	7819	9167	9618	8408	7942	7789	8322	8543	7568	7715	8783	7208	9278	7118	8552	,	9110	6142	7689	7902	7596
(NB)		DM	%		Н	55.1	8.0	52.4	ο.	9	32.8	œ.	ζ.	27.4	ö	7 .	ω.	о О	21.7	4.	37.9	2		<u>.</u>	9	38.8	•	•
Test 599 (Stand	Count	Mean	(cont.)	ė.	ø.	16.3	9	9	9	16.3	9	5.	15.7	9			•	16.0	•	15.7	5.		و	ė.	16.3	ė.	9
Η		%Bolt	8/26	PX	38.9		2.1			37.7	0.0	32.4	0.0	40.4	20.7	92.0	0.0	69.7	2.0	51.8	76.5	œ.	R780/2			6.3		œ.
		Variety		R880-# = RZM R780	1	4 -	I S	9 1	- 7	80 I	6 I	-10	-11	-12	-13	-14	-15	-16	-17	-18	-19	-20	"]		- 2	m I	4 -	ا ج

EVALUATION OF PAIRCROSSES (FULL SIBS) OF C78 & C80, SALINAS, CA., 1999

(TESTS 599, 1599, 4599)

i	Te	- 1	(NB)		Test	1599	(VY)	i		Test	4599 ((Rzm)	
%Bolt		Stand Count	MQ	Sugar Yield	Sucrose	RJAP	Beets/ 100'	Score	Sugar Yield	Sucrose	RJAP	RZM Resist	Root Rot
8/26	I	Mean	o,e	1bs	∞	o%	No.	Mean	1bs	æ	₩	1	∞
R780/2		PX (cont.	<u></u>										
	l	16.3	32.8	7354	6.2	4.	121	4.9	ın	6.1	4.	91.7	6.7
49.5		•	17.8	8477	6.3	ω.	Э		\sim	7.0	4.	0	8
45.0		ė.	26.7	8527	6.4	ω.	ω		O)	7.3	ъ.	8	•
36.8		13.7	•	7765	16.63	83.2	118	5.2	8441	17.90	84.3	83.5	9.5
2		•	17.0	6148	6.3	Ή.	Н	•	യ	6.5	4	ο 0	•
0.0		15.0	44.4	8024	6.6	æ.	4		o	7.2	Ŋ.	m.	0
R780-	-45	5 PX											
12.0	1	16.7	31.9	48	5	84.4	4	•	8552	ø.	m.		8
22.6		16.0	51.5	9884	16.07	83.4	152	4.8	6770	16.60	82.6	82.5	13.0
•		15.7	•	01	9	2	9	•	8655	7.	8	о	•
,		7	7 7	r	u	ď	r		_	u	u	0	
•		•) α) (1	•	10) (•
			٠ ر	8049	15 73	, w	130	. 4	7811	15.03	יי ס מ	0.7.0	ο α
•		•) i)		•	` (•	4 1	;	,	:	
2.4				_	6.0	⊢	4	•	8155	9	, ,	m	14.4
79.2		•	ö	8178	ъ.	ω.	ന	•	91	6.5	ά.	5	•
•		<u>ي</u>	0	9507	ė.	4.	ω	•	05	6.8	4.	ω.	•
4.4		15.3	16.9	8214	16.00	82.6	142	4.7	7999	17.33	84.3	95.0	4.2
10.9		5	<u>و</u>	6102	IJ.	4.	m	•	31	6.7	m.	5.	•
0.0		15.7	•	92	6.1	8	\vdash	•	61	6.7	w.	Η.	•
•		•	32.3	7582	16.37	83.0	121	5.0	7952	16.73	83.4	72.5	2.1
5.1		14.0	38.8	22	6.5	8	\vdash	•	31	6.6	H.	8	•
•		•	38.0										
									8157	16.97	84.6	82.8	15.6

EVALUATION OF PAIRCROSSES (FULL SIBS) OF C78 & C80, SALINAS, CA., 1999

(TESTS 599, 1599, 4599)

	Tes	Test 599 (NB)			Test	Test 1599 (VY)	Y)			Test	Test 4599 (Rzm)	Rzm)	
		Stand		Sugar		_	Beets/	ΔX	Sugar			RZM	Root
	%Bolt	Count	DM	Yield	Sucrose	RJAP	1001	Score	Yield	Sucrose	RJAP	Resist	Rot
	8/26	Mean	%	1bs	%	œ۱	No.	Mean	1bs	o,e	% I	 %∣	ا %
~1	R870-# = RZM R770 PX (cont.)	sont.)											
	29.6	15.7	49.0						7723	16.33	85.9	69.5	35.8
	46.4	16.3	26.3						9301	17.67	83.1	90.5	3.5
	33.4	16.0	29.9						7154	17.13	82.5	86.7	50.2
	13.6	17.3	47.9						8134	16.87	83.9	100.0	24.2
	27.7	15.8	37.1	7926.	7926.3 16.17	83.4	141.7	4. 9.	7851.3	7851.3 16.69	83.7	85.6	11.6
	18.8	1.9	28.7	1758.9	9 0.84	2.5	26.5	9.0	1823.9	0.79	3.2	13.4	26.1
	42.0	7.3	47.8	13.7	7 3.20	1.8	11.6	7.5	14.4	2.94	2.4	9.7	138.9
	14.5**	×*6.8	3.5**	ώ.	3.7**2.37**	1.8**	1.8**	* 3.7**	2.1	2.1**3.57**		1.3NS 5.9**	1.3NS

See Tests 1599 and 4599 for performance under virus yellows and rhizomania. TEST 599 NOTES:

TEST 1599 NOTES: See tests 599 and 4599 for companion tests under bolting and rhizomania conditions. Inoculated with VY (BYV-BWYV-BChV).

TEST 4599 NOTES: Harvested by hand with 5% tare. Lifted roots counted (harvest count), rhizomania susceptible roots counted (% RZM resistant), and root rot counted (Sclerotium rolfsii). Total plot weighed but only nonrotted roots used in sugar sample. Also see results from bolting and virus yellows trials.

EVALUATION OF MULTIGERM S1 PROGENY LINES FROM F1 POPULATION HYBRIDS, SALINAS, CA., 1999

(TESTS 699, 1699, 4699)

	Root	Rot	%		4.3	20.4	2.2	16.2		14.6	ო				7		14.7		7.0	•			7.8		8		4.4	•		•
(Rzm)	RZM	Resist	o⁄o I		5	84.3	8	e.	,	76.9	ω ω			9	10.5		m.		•	•	•	•	59.3	•	•	•		9	71.3	7 .
4699 (R		RJAP	%		ς.	80.1	N.	Η.	(83.7	85.1			4.	81.8	. (83.2		•	80.0	•		81.8					4.	81.6	0
Test		Sucrose	& 		6.2	16.47	7.1	6.4	,	16.43	6.5			7.6	14.27	1 T	5.1		ė.	ъ.	ø.	6.	16.00	9	ъ.	9	9	9	16.47	9
	Sugar		1bs		86	7296	40	55	(7905	25			ന	5134) (∞		6758	6236	7671	6394	8039	0969	7185	8124	6626	σ	6246	8155
	ΛĀ	Score	Mean			3.6	4.8	•		•	4.3	4.3	•						•	5.0	•	•	4.2	4.1	•	4.2	•	•	4.3	•
(2	Beets/	1001	No.		161	152	139	145	,	136	148	170	164						179	170	148	136	155	130	127	136	4	ъ	142	Ŋ
(VV) 6691	Н	RJAP	₩		Ή.	6.08	е Э	2	,	0	ъ.	84.4	o.						80.2	•	•	•	79.4	•				ζ.	81.4	2
Test 1699		Sucrose	≫ I		4.9	15.27	6.5	5.6	L	υ α	6.2	15.87	ω.					-5	9.	Τ.	ω.	5.8	15.03	۲.	ω.	5.0	5.2	5.5	15.60	5.3
	Sugar	Yield	1bs		7377	7627	7450	7258	0	8307	10418	7035	3649					x C76-89-5	7240	7181	6645	6097	10101	8249	4722	8011	6752	8625	7606	7964
(NB)		DM	or		48.0	•	6.1	•	(עכ	39.0	24.1	9					913-70		26.2			•		36.6	•	65.2	٠	•	42.0
669	Stand	Count	Mean		17.3	15.0	16.7		L	15.0	ю		ė.					13⊗ = C	6.	16.3	7.	9	9	15.0	16.0	9	16.3	7.	•	16.7
Test		%Bolt	8/26		21.2	39.4	36.4	•	,	4.	4	15.9	8					R776-89-5H13⊗	•	•	•	•	16.3	•	•	21.8		7.7	•	6.1
		Variety		Checks	8913-70	8918-12	R886-89-5NB	8935 (Iso)	(8939	X869	97-C37	97-SP22-0	B4776R	US H11		TH SO	8935-# = RZM F	ı	- 2	m I	۱ 4	ا ت	9 1	- 7	80 I	O I	-10	-11	-12

(TESTS 699, 1699, 4699) (cont.)

	Root	Rot	%		0.0	0.0	8.3	•	8.3	33.8	0.0	14.7	2.1	•	9.5	•	•	0.0	2.1	•	3.9		30.3	2.4	•		, r	
zm)	RZM	Resist	o o		6.97	91.5	8		•	74.5		82.4	9	ъ Э	83.6	0	89.4	94.4			97.9		•	74.0	•	0	0.10	_
Test 4699 (Rzm)		RJAP	%		81.1	0	79.5		81.4	•	82.6		•	$\ddot{\mathbf{H}}$	83.4	6	•	82.2	•	•	82.9			81.9	ω.		2.10	-
Test	i) -	Sucrose	%		16.83	16.63	15.70	5	7.	16.37	°.	15.67	6.4	6.	16.53	6.8	7.8	16.90	6.8	ю.	15.70		6.5	15.07	9.9	2	16.27	
	Sugar	Yield	1bs		7522	6755	7261	0209	35	5856	0099	41	\vdash	\mathbf{H}	8109	9	7221	6270	7291	5979	8430		2	6756	m	8	8270	ก
	ΛX	Score	Mean		4.9	5.0	5.1	4.2	4.7	4.2	4.3	4.2	4.5	4.3	4.2	4.3	4.4	4.8	4.6	4.2	4.3		4.6	4.6	4.4) C	٠
Y)	Beets/	1001	No.		145	148	148	133		4	133	4	~	ω	145	4	161	136	142		139		m	133	က	-	1 20	7
1699 (VY)		RJAP	∞		•	82.1	•	4.		•	81.9	•	77.2	8	81.5	m.	H.	83.1	÷.	æ.	82.6		•	81.3	ω.	0		į
Test 1		Sucrose	અ	\sim	6.0	15.93	4.7	4.8	5.9	5.2	15.47	4.7	5.3	5.7	15.17	5.9	6.8	15.40	5.9	5.2	15.23		6.4	13.37	5.3	8	1 1 1 0 0 1 1 1 1 0 0 1 1 1 1 1 1 1 1 1	٦. ۲
	Sugar	Yield	1bs	-89	8164	7138	7229	7817	6891	8571	ന	⊣	05	19	9327	71	6319	7248	7055	7122	6122		8105	6212	8500	6407	70.07	1223
(NB)		DΜ	₩	C913-70		19.2	•				11.3	•	4.	8	34.6	ö	•	0.0	•	42.2	44.0		4.	32.9	ъ.			Ÿ
Test 699 (1	Stand	Count	Mean	11		15.7	5	5	9	16.3	15.0		9	•	15.3	5	•	13.7	•	14.3	16.7	H31⊗	16.7	15.3	ъ.	v		ė
Tes		%Bolt	8/26	R776-89-5H13®	11.1	66.3	•	ъ.		•	37.9	•	51.4	9.6	•		14.4	7	6.5	27.8	34.5	R776-89-5H31⊗		82.1	•			٠
		Variety		935-# = RZM	8935 -13	-14	-15	-16	-17	-18	-19	-20	-21	-22	-23	-24	-25	-26	-27	-28	-29	8936-# = RZM	936 - 1	- 2	ا ع	- 4	י על ו	

EVALUATION OF MULTIGERM S1 PROGENY LINES FROM F1 POPULATION HYBRIDS, SALINAS, CA., 1999

(TESTS 699, 1699, 4699) (cont.)

	Root	4	I		12.5	2	•	0.0	•	•	5.8		23.1		6.7	4.5	0.0	0.0					10.0		•
(Rzm)	RZM	וו	1		81.3	•	•	90.3	•	93.6	9		70.0	•	w.	4.	75.6	4.			7.	Б.	0.09	0	9
4699 (1	Q.T.A.	*	1		90.8	m.	•	72.7	•	4	m	5.	82.0	•	•	Э.	80.1	ω.			o.	e.	83.2	6	2
Test	Sucross	% 01010 01010 01010	I		16.87	7.0	7.8	15.83	7.0	6.4	6.7	6.2	17.23	6.1	6.7	6.0	17.60	6.7			6.5	5.3	16.87	7.1	6.7
	Sugar	1bs			7478	8903	7536	6171	8316	8706	7374	4392	6807	8454	7337	7147	6663	9018			6665	5658	6450	5650	6201
	VY	Mean			4.3	4.2	4.8	4.9	5.2	•	•	•	9. ₉	•	•	•	4.7	•	ი. გ		•	•	5.6	•	•
Y)	Beets/	No.			130	N	⊣	142	$^{\circ}$	124		ന	142	ന	7	ന	142	\vdash	133		133	118	130	118	145
1699 (VY)	041.0	% %	·I		82.3	m.	•	80.6		82.9		•	81.5	•	Η.	8	9.62	m.	83.3		•	•	84.3	•	•
Test	0000000	Sacrose *	·I		14.90	6.2	8.0	15.67	6.7	16.30	ъ.	5.5	16.00	16.03	6.9	4.0	16.50	4.9	16.30		6.1	5.3	16.23	9.9	6.1
	Sugar	1bs			8226	10205	6543	7086	7253	7373	6846	6052	7822	7180	7173	6300	7976	8935	6470		6978	4598	5704	6901	6233
(NB)	2	*	· i	(cont.)	21.4	2.4	11.4	21.1	55.7	53.2	48.5	46.4	14.6	21.4	•	•	100.0	•	13.5		ė.	•	0.0	•	5.0
669	Stand	Mean			14.3		14.7	5	15.7	16.3	13.7	14.7	•	15.3	13.3	•	•	14.0	15.0		•	کا	15.7	کا	m.
Test	9 1 1	8/26	٠l	R776-89-5H31⊗	67.2	79.8	4.8	62.9	19.0	0.0	32.6	14.2	9	59.5	20.0	90.5	31.1	54.4	10.7	R776-89-5H11⊗	3.	•	42.9	•	37.0
	170	variety		8936-# = RZM	8936 - 6	7 -	80 1	6 I	-10	-11	-12	-13	-14	-15	-16	-17	-18	-19	-20	8937-# = R77		- 2	۳ ۱	4 -	ហ

(TESTS 699, 1699, 4699) (cont.)

(Rzm)	RZM Root	Resist Rot	o,e o,e		2.	8.0 29.	2	.3 0.	1.8 12.	7.1	0.0	1 4.	13.	5.6 23.	7.9 11.	.6 4	.9 10.	4.1 0.		4.9 19.				.8	45.9 14.3	o.	21.5 21.6	9.1 140.
4699 (R		RJAP	o⊳ I		8.7	3.1		1.2	2.2	α) r	ım		4.7	1.3		2.3	9.7	m.	4.6					81.0	81.4	.5	
Test		Sucrose	o⊱l		6.7	6.5	16.20	6.8	4.5	Δ	. 6	16.17	4.2	3.2	16.50	5.3	6.2	ъ.	16.67	5				4.	14.97	16.31	1.20	ഹ
	Sugar		1bs		7025	7026	87	7261	7199	6759	6764	7751	~ ~	78	സ	6229	55	5637	6673	8009				7018	5728	12.	1832.3	
	ΔŽ	Score	Mean		4.4	•	4.5	•	•		•		•	•	5.0	•	4.6	4.9	4.6	4.5	9. 8	4.1				4.6	0.5	•
Y)	Beets/	1001	No.		ന	2	127	Ω	2	-	٠.	121	-	4	133	m	⊢	Н	133	4	⊢	124				137.1	25.0	_
1699 (VY)		RJAP	o(0		ک	4.	81.4	ŀ.	т М	0		81.3	84.8	5	83.9	H.	ω.	9	83.1	ω.	Ή.	86.1				82.2	3.0	
Test		Sucrose	o/0		5.2	5.9			4.7		1 9	15.57	4.0	3.9	Н	16.63		5.0	16.47	5.8	5.5	15.20				15.5	6 1.00	ď
	Sugar	Yield	1bs		7471	ဖ	6619	6114	0	Ľ) LC	7158	4	6336	13	8636	8746	α	6453	α	7433	9168				20	1987.6	_
(NB)		DM	o⊱1		•	49.1	64.2	15.4	41.1	ע		11.3	16.7	37.5	0	20.2	21.9	4.8	51.4	7.2				ω.	48.1	31.0	9	ر 1
669	Stand	Count	Mean		14.7	4.	15.7	5.	رى	۷	13.7	•	13.0	4.	14.0	•	12.3	4	15.0	5.				•	15.3	ω.	20.9	α
Test		%Bolt	8/26	m I	29.5	•	2.2	•	57.5	Ľ	0		45.7	38.4	9	34.4	ი	30.3	29.0	•			CR812-#		53.4	15.3	2.0	0,88
		Variety	=	39-# = RZM	8939 - 1	- 2	۳ ۱	4	ı S	ı	- 7	- co I	თ I	-10	-11	-12	-13	-14	-15	-16	-17	-18	CR811-# & CR8	CR811-1	CR812-1	Mean	\sim	C. ⟨%)

EVALUATION OF MULTIGERM S1 PROGENY LINES FROM F1 POPULATION HYBRIDS, SALINAS, CA., 1999

(TESTS 699, 1699, 4699)

	Root	Rot	o,0
Rzm)	RZM	Resist	%
Test 4699 (Rzm)		RJAP	æI
Test		Sucrose	o o
	Sugar	Yield	1bs
	$\Lambda\Lambda$	Score	Mean
VY)	Beets/	1001	
1699 (VY)		RJAP	%
Test 1699		Sucrose	અ∘
9	Sugar	Yield	lbs
(B)		DM	æ
st 699 (1	Stand	Count	Mean
Te		%Bolt	8/26
		Variety	

TEST 699 NOTES: See Tests 1699 and 4699 for performance under virus yellows and rhizomania. See notes for Test 299.

 $RZM R776-89-5H138 = 6913-70aa \times R576-89-5$ RZM R776-89-5H318 = 6931aa x R576-89-5 8935-# =

 $R776-89-5H110 = 5911-4aa \times R576-89-5$

= #-9868

RZM Y769H31 \otimes = 6931aa x Y669 8937-# = = #-6868

R576-89-5 = C176-89-5 = Inc. of full sib family selected from popn-76-89. C913-70 = C913-70

6931 = MM,S^f, Aa, Rz population

X669 = C69

5911-4 = C911-4

See tests 699 and 4699 for companion tests under bolting and rhizomania conditions. Inoculated with VY (BYV-BWYV-BChV). TEST 1699 NOTES:

Lifted roots counted (harvest count), rhizomania susceptible TEST 4699 NOTES: Harvested by hand with 5% tare. Lifted roots counted (harvest count), rhizomania sustroots counted (% RZM resistant), and root rot counted (S. rolfsii). All roots in plot weighed but only nonrotted roots included in sugar sample. Also see results from bolting and virus yellows trials.

(TESTS 799, 1799, 4799)

	Root	Rot	ا %		Η.	9	9		8	8.6	i.	5		9	ന	•	ف		1.7			•	Η.	3.9	•	80	6.1	9	2
m)	RZM	Resist	o%		5.5	ω.	8.3	5.	6.	9.4	•	٠ و		9.7	7.2 1	1.	m.	6.0	2.2 2			m.	8	σ	9	ω.	5.7	ij	5.6
4799 (Rzm)		RJAP R	1%		4.0 6	5.9 7	5.9 7	4.7 7	3.8 8	2.9 5	3.8	5.4 1		4.1 7	1.6 8	9.4 7	3.4 8	7.3 8	7.3 8			4.7 6	9.5 7	6.1 8	5.0 7	4.0 8	1.8 7	2.3 7	2.8 7
Test 47		- {								.07				80	.37 8	57	43	80	.73 8			87	53	.33 8	00	17	.40 8	93	43
H		Sucrose	6 0		\mathbf{c}	9	9	9	2	15	ന	ന		15	16	16	15	7	15			15	15	15	16	15	15	15	15
	Sugar	Yield	1bs		9	7	87	64	83	6786	9	87		8091	7304	7119	1906	74				m	7136	7790	0699	-	7736	0	m
	ΛĀ	Score	Mean											•	5.8	•				5.7		•	•	5.7	•		5.7	•	•
(۲۸)	Beets/	1001	No.											152	152	158	136	142	158	161	155	m	ω	152	4	4	142	$^{\circ}$	$^{\circ}$
1799 (1		RJAP	o(0											2	82.2	9	4.	2	5	82.4	9	4.	4.	85.9	۲.	8	81.1	ë.	2
Test		Sucrose	æ1											6.2	16.30	6.1	5.2	6.5	6.6	17.30	5.6	6.4	6.0	15.83	6.0	6.2	15.83	6.5	6.2
	Sugar	Yield	1bs											8099	6629	5501	7777	8984	87	7927	39	54	26	6046	22	26	6333	57	60
(NB)		DM	₩	σ,	23.6	,			42.8					40.3	78.3	2.2	54.7	9	ك	55.1	2		ω.	64.7	4.		14.6	•	25.6
Test 799 (Stand	Count	Mean	ý	17.0				16.7				118	16.7	•	15.0	9	15.7	9		•	14.7	5	16.0	5.		15.3		5.
E		%Bolt	8/26		33.3	,			32.6				ER-8S 6931®	11.9	2.2	0.0	•	0.0		•	•	38.9	2	45.3	•	0.0	2.1	•	60.1
		Variety		Checks 7933	8931	8931	8931	8931	8926 (Iso)	· w	S H1		8931-# = RZM-ER-%	8931 - 1	ı	რ I	- 4	l L		- 7	80 I	თ I	-10	-11	-12	-13	-14	-15	-16

EVALUATION OF MULTIGERM S1 PROGENY LINES FROM RANDOM-MATED POPULATIONS, SALINAS, CA., 1999

(TESTS 799, 1799, 4799) (cont.)

1	Root	Rot	%		9.5	0.0			15.4	9			22.6		2.4	12.1	4.4			_	26.7		0.0	11.7	0.0	0.0	20.6	
(Rzm)	RZM	Resist	%		87.3	71.6				94.2		5	71.0	8	91.5	8.76	87.3		Ų	0.0/			94.9	87.6	82.8	0.00.	66.4	
4799 (R		RJAP	e/e		4.3				4.	81.1		6.	86.0	ω.	4.	84.9	e.			43.1	82.7		74.7	83.4	0	m.	81.9	
Test		Sucrose	%		15.53	14.03			5.3	16.20		14.87	15.07	9	S.	15.03	9		L	ů.	14.00		17.30	16.07	16.10	15.60	15.53	,
	Sugar	Yield	1bs		6762	7266			5917	6484		7548	6972	8327	6638	9177	6801		000	909/	7273		5254	7466	7126	8673	6679	
	Λλ	Score	Mean		4.5	5.8		5.2	5.8	5.0		5.1	4.9							ი.ი	•	4.5						
<u>۲</u>)	Beets/	1001	No.		127	130		130	127	142		124	139						1	145	115	152						
1799 (VY)		RJAP	₩			83.2		83.6		84.3		84.2	ω.								80.0	81.2						
Test		Sucrose	o⊱		16.07	15.30		16.77	15.77	16.27		16.10	15.67						L	•	15.20	16.37						
	Sugar	Yield	1bs		7728	6450		7930	6524	6778		7079	8757							6779	3989	6587						
(NB)	1	DM	æΙ	·:	56.6	41.3	8	28.6	8	15.6		•	37.7	2.	62.2	4.4	32.0			V	9.69	10.4	60.8	71.0	59.5	•		
Test 799 (Stand	Count	Mean	1⊗ (cont	15.0	14.7	10.7	14.0	4			14.3	14.7	•	•	15.3	e		L	ر د	15.3	16.7	13.0		14.0	4		
Tes		%Bolt	8/26	-ER-%S 6931⊗	٠.	90.5	•	•	14.0		,	6	28.9	9	48.2	8.6	25.3	70268		'n.	23.9	76.1	15.7		26.2	22.4		
		Variety		8931-# = RZM-ER-	8931 -17	-18	-19	-20	-21	-22		-23	-24	-25	-26	-27	-28	1 7 7 7	1777	8926 - I	- 2	m I	- 4	۱ ک	9 -	- 7	80	

(TESTS 799, 1799, 4799) (cont.)

	Root	Rot	% I		14.1	6.7	24.1	33.0	38.8	28.6			5.9	10.1			13.8	4.6	3.4	2.5**
(Zm)	RZM	Resist	%		60.4	41.5	5.9	24.4	2.4	59.0			71.7	70.6			70.7	22.8	19.9	**6.8
4799 (Rzm)		RJAP	o(P		81.7	77.0	83.0	85.5	81.5	86.3			79.0	83.6			83.6	4.6	3.4	2.5**
Test		Sucrose	%1		16.17	16.83	14.27	15.50	13.60	16.30	15.53	15.77	15.47	14.30			15.55	0.93	3.67	1** 8.30**
	Sugar		lbs		6495	5802	4669	5298	3919	6531	6800	6899	5068	5227			6923.3	1731.7	15.4	4.1
	ΜĀ	Score	Mean		5.6			6.3	6.0		5.8				5.2	4.4	5 .5	0.5	5.7	* 7.7**
(VY)	Beets/	1001	No.		121			103	127		130				145	145	139.0	25.3	11.2	2.2*
1799 (V		RJAP	o40		80.7			82.2	85.3		83.0				84.1	82.6	82.8	2.5	1.8	3.1**
Test		Sucrose	o%	Res.)	16.57			16.43	16.03		15.87				16.17	15.63	416.08	5 0.57	5 2.17	6**5.95**
	Sugar	Yield 8	1bs	aphid	5826			6516	5308		6138				7225	5501	6800.4	1830.	16.	2.
(E)		ΩM	%	338 (Root	70.8	44.4	54.6	47.9	35.4	41.7	38.2	38.0	4.4	23.5			41.1	31.7	47.5	3.8**
Test 799 (NB)	Stand	Count	Mean	7227,79338	15.0	15.0	13.3	15.0	15.3	15.0	16.3	16.7	15.7	15.7			29.9	19.0	39.2	11.4**
Tes		%Bolt	8/26	RZM 7221,7225,7227	42.5	22.2	45.2	33.4	11.0	9.1	22.5	0.0	37.4	51.0			15.1	1.7	7.0	3.8**
		Variety		8933-# = RZM	8933 - 1	- 2	m I	-21	-22	-23	-31	-32	-33	-34	-41	-42	Mean	LSD (.05)	C.V. (%)	F value

See Tests 1799 and 4799 for performance under virus yellows and rhizomania. TEST 799 NOTES:

TEST 1799 NOTES: See tests 799 and 4799 for companion tests under bolting and rhizomania conditions. Inoculated with VY (BYV-BWYV-BChV).

TEST 4799 NOTES: Harvested by hand with 5% tare. Lifted roots counted (harvest count), rhizomania susceptible roots counted (% RZM resistant), and root rot counted (Sclerotium rolfsii). All roots in plot weighed but only roots without rot included in sugar sample. Also see results from bolting and virus yellows trials.

EVALUATION OF MONOGERM S_1 PROGENY LINES FROM RANDOM-MATED POPULATIONS, SALINAS, CA., 1999

(TESTS 899, 1899, 4899)

()		Kesist Kot	19.	82.1 30.5			15.0 57.5			15.	65.6 26.4		,	3 34.	87.0 30.0	2 15.	9 49.	18	3 28	57.0 28.9	3 26	
4899 (Rzm)		KJAP Ke		81.2 8			82.3 4		7	ω.	82.7	٦.	,	٥.	83.3	.7	6.	0.	.7		ω.	
Test	,	Sucrose	15.37	ß			14.47		14.87	15.30	15.57	14.67	Ł		16.00			14.73	16.27	15.60	15.83	
	Sugar	ibs	6132	4767			9089		3622	5430	5844	6193	i C	2847	3925	4845	5466	5381	5019	5341	5157	
	ΔĀ	Mean	e. 9	5.6		5.6	•												5.6		5.5	
7Y.)	Beets/	No.1	158	161		145	91												109		136	
Test 1899 (VY)		KOAP *I	m	81.7		83.9	œ.												80.7		77.8	
Test		Sucrose	16.13	15.50		15.60	14.60												15.97		15.80	
	Sugar	Yield	8051	7102	~ l	5163	6196												3255		3778	
(NB)		Σ % Ι	2.0	2.2	testcrosses	2.2	5.1		11.1	22.3	8.3	14.1		Ω Ω	2.5	35.7	16.2	4.4	16.1	12.9	15.2	
Test 899	Stand	Count	16.3		(T-0 tes	14.7	13.3		14.7	13.3	16.0	16.7	ı	٠	٠	13.7	14.3	13.7	14.7	15.7	15.3	
Ĕ		%Bolt 8/26	0 05		7838mm⊗	13.7	30.0	RZM 7835mm⊗	13.9	42.3	29.2	45.8		8.0/	0.0	0.0	7.0	14.9	0.0	10.8	2.1	
		Variety	Checks	8838m	8838-# = RZM	8838 - 5	9 1	11	8835 - 3	- 4	ا 5	9 -	ţ	/ -	ი 1	-10	-12	-14	-15	-16	-17	

EVALUATION OF MONOGERM S1 PROGENY LINES FROM RANDOM-MATED POPULATIONS, SALINAS, CA., 1999

(TESTS 899, 1899, 4899)

(cont.)

EVALUATION OF MONOGERM S1 PROGENY LINES FROM RANDOM-MATED POPULATIONS, SALINAS, CA., 1999

(TESTS 899, 1899, 4899)

(cont.)

	Root	₩		12.5	17.4		30.7	5.6	42.4		20.6	8.6	10.6	0.0	!		7.8	0.0	7.8	0.0	1.6	12.5	4
(Rzm)	RZM Resist	o%		83.5	94.1		76.4	84.4	81.4		٠	٠	74.7	76.3	 		74.1	78.8	69.8	39.8	83.8	ω	2
4899 (R	RJAP	. ₩ I		83.1	4.		82.4	79.5	80.4		82.2	٠	80.6	81.5	1 1 1			82.1	83.6	84.5	•	83.3	•
Test	Sucrose	o, 0		16.07			16.03	•	15.87		Ď.	<u>ب</u>	15.77	16.17	! ! !		•	16.03	15.97	15.40	16.10	•	9
į	Sugar Yield	1bs		7184	4721		4666	4381	9909	- 1	8/19	5565	5448	6740	 		89	5979	4917	4880	6272	ന	4829
	VY Score	Mean					6.0								1 1 1 1		•	6.3	5.7	5.8	5.8	5.8	5.3
Y)	Beets/ 100'	No.					139								1 1 1		130	121	130	139	158	127	142
1899 (VY)	RJAP	o⊳ I					81.3								1 1 1		82.1	81.4	80.7	82.5			6
Test	Sucrose	o⊱					16.27								1 1 1 1		•	16.00	•	16.07	15.50	6.4	4.
-3	Sugar Yield	1bs					4886								1 1 1		6029	3829	3902	5551	3940	7142	3414
(NB)	MO	æ		37.4	4.2		17.4	29.7	14.7		6.7	4.8	19.7	0.0	1 1 1		2.1	0.0	0.0	4.2	10.7	6.5	0.0
Test 899 (Stand	Mean	⊗	14.0	14.3	⊗.	15.0	16.7	15.7		15.3	14.0	14.0	15.0	1 1 1	6828mm⊗	16.0	15.3	15.7	15.7	16.0	4	14.7
Ξ	%Bolt	8/26	RZM 7835H69mm⊗	0.6	2.1	RZM 7835H87mm⊗	51.0	0.0	57.3		23.9	25.9	10.8	97.8	1 1 1	RZM-ER-8S 682	56.3	41.9	27.1	4.2	49.2	0.0	13.5
	Varietv		11	8835 -61	-62	II	8835 -71	-72	-73	i	-74	-75	-76	-77	1 1 1 1 1	11	8828 - 1	- 2	e I	- 4	ى ا	9 -	- 7

(TESTS 899, 1899, 4899)

(cont.)

	Root	Rot	oke		5.9	6.3	0.0	4	23.7	6		7.4	0.0	22.1	0.0			30.9	39.8		5.	12.4	9	;
(Rzm)	RZM	Resist	o⊱l		95.5	9.68	8	9.62	ά.	64.0				95.6				93.0	74.0		•	86.9		
4899 (F		RJAP	%		82.8			82.6	81.0	83.9			•	79.2			•	83.4	81.5			80.8		
Test		Sucrose	æ		15.50	15.33	14.70	15.70	15.87	14.40		16.73	17.20	15.67	15.77		•	16.70	15.90		16.17	15.07	16.50	16.70
	Sugar	Yield	lbs		5815	5509	5347	5564	4635	3707		6186	5853	5950	5637		7857	5535	5117		6406	5998	7126	6329
	ΛX	Score	Mean		•		6.4	6.7	5.7			6.5									5.8	5.6	6.4	5.4
X)	Beets/	1001	No.		139	179	152	148	161			145									139	121	133	148
1899 (VY)		RJAP	oko		•	•	•	84.6	•			82.5										80.0		
Test		Sucrose	₩		4	r	E	15.33	2			15.40									9	14.93	Ŋ	9
	Sugar		1bs		4576	4794	4594	4214	6105			3419									6983	5305	6556	5397
(NB)		DM	% ∣		11.4	7.9	6.1	0.0	34.7	2.0		4.0	19.9	6.4	17.6		6.8	8.1	4.2		30.3	•	4.9	10.7
Test 899 (Stand	Count	Mean	6833mm⊗	15.7	17.3	16.0	15.3	16.7	15.3	6833%mm⊗	16.0	15.3	15.0	9.6	68348mm⊗	15.3	16.0	15.7	6836mm⊗	15.3		13.7	15.3
Te		%Bolt	8/26	RZM-ER-%S 683	40.1	44.9	41.4	22.3	34.4	60.4	RZM-ER-8S 683	68.3	34.7	15.8	22.0	RZM-ER-8S 683	22.0	3.9	89.5	RZM-ER-%S 683	0.0			8.9
		Variety		11	8833 - 1	- 2	ო 	- 4	۱ 5	9 1	II	8833 11	-12	-13	-14	8834-# = RZM-	8834 - 1	- 2	რ I	!!	8836 - 1	- 2	ო 1	4

EVALUATION OF MONOGERM S1 PROGENY LINES FROM RANDOM-MATED POPULATIONS, SALINAS, CA., 1999

(TESTS 899, 1899, 4899)

(cont.)

	Root	Rot	% I			13.9	ω.			5.6	•		72.0	0.0	13.5		26.9	15.7	6	36.0	41.4	•	14.6	23.7	36.8
(Rzm)	RZM	Resist	% I			35.7	9.68		0	81.8	m.	(76.9	ق		9	90.7			ъ.	87.4	÷.	ω.	77.8
4899 (R		RJAP	o%			\vdash	•		æ.	81.2	2	('n	81.5	•		84.1	82.4	77.3	4	ت	82.1	ë.		83.5
Test		Sucrose	%			16.30	6.9		5.5	15.73	5.7	1	۲. ۲	16.23	6.9		Ŋ		ت	14.77	ъ.	15.60	IJ.		16.70
	Sugar	Yield	lbs			5583	7775		8	5291	4		4 R	6116	77		5525	91	36	m	N	6622	0	82	7992
	ΔŽ	Score	Mean						•	6.1	•		ა.ა	5.8	5.8		5.8	6.5	6.0		•	5.8	•	•	5.8
X)	Beets/	1001	No.						$^{\circ}$	155	4		121	127	136		$^{\circ}$	139	ന	127	142	136	152	വ	152
1899 (VY)		RJAP	%						æ.	81.8	2	1	٠	82.0	81.5		Η.	81.4	0	•	•	82.2	•	•	81.7
Test		Sucrose	%						5.9	15.73	6.4	,	9.5	16.23	6.7		5.8	15.10	6.4	5.5	5.5	15.73	5.7		16.63
	Sugar	Yield	1bs						6894	5767	4948	1	7438	6601	6119		4878	3385	4507	3326	5947	6557	6582	5371	6617
(NB)		DM	o⊳ I	ont.)	21.6	8.3	19.1		0.9	0.0	0.0	0	32.9	2.2	0.0		4.2	4.4	3.9	0.0	0.0	2.1	3.9	6.8	15.4
Test 899 (Stand	Count	Mean	6836mm⊗ (cont.)	10.0	15.3	14.0	6837mm⊗	16.7	16.7	16.0	1	•	14.7	15.0	⊗	15.3	14.7	16.7	15.3	15.7	16.0	16.3	15.0	15.3
Te		%Bolt	8/26	RZM-ER-8S 683	0.0	0.0	0.0	RZM-ER-%S 683	42.3	2.0	85.4			54.8	0.0	RZM-8S 6808mm⊗	48.5	18.3	0.0	35.4	0.0	31.5	47.2	44.4	15.3
		Variety		11	8836 - 5	9 -	- 7	11	8837 - 1	- 2	m I	•	- 4	ا ت	9 -	8808-# = RZM	8808 - 1	- 2	ю 1	- 4	ı S	9 -	- 7	ω 1	თ I

(TESTS 899, 1899, 4899)

(cont.)

	Root	Rot	%		5.0	36.7				23.9		2.0		0.0		C	14.8) •	•	22.9	•	5	3.6		8.4	6.9	m.
~	l	Resist	∞ા		.7	86.4 3		७.	0.	ο.		8.0		9.9		v.	0	`	.2	4.		9.	0.0		0	.8	.7 3
4899 (Rzm)	1	RJAP Re	₩		ო.	7		٦.	.7 1	ω.	m.	.2		8		7	68		9.	0.			.6 20		80	.3 71	
489		- 1			84	വ		82	83	82	82	84		81		83	2 6	5	82	82	82	0	85		80	83	78
Test		Sucrose	o(P		16.20	14.93		15.93	15.97	14.90	16.50	16.00		15.60		15.37	15.87		15.67	15.77	15.43	4.	14.27		15.47	15.93	15.77
	Sugar	Yield	1bs		5240	6316		4969	5739	4316	6434	7209		4876		4906	5169) 	6597	5963	3920	6716	3982		4319	5843	3571
	ΔĀ	Score	Mean		5.4	5.8		•	•	5.3	•																
(X)	Beets/	100'	No.		121	130	,	121	106	145	145																
Test 1899 (VY)		RJAP	æ i		82.2	84.2	,	N.	4.	84.6	9																
Test		Sucrose	ονi		15.80	15.60	,	16.17	15.67	15.53	16.60																
	Sugar	Yield	lbs		4099	5151	(8929	2988	2451	3886																
(NB)	i	Σ Ω	ον I	·	2.0	0.0		4.5	9.7	10.7	4.2	17.8	4.8	15.0	14.4	18.3	23.5)	6.7	12.5					3.7	2.0	6.8
Test 899	Stand	Count	Mean	<u> ⊗</u> (cont.)	16.7	13.7		ď.	•	5	15.3	13.3	5.3	13.0	14.0	13.7	ري	1	15.7	15.7				8	16.3		15.7
Te		%Bolt	8/26	RZM-%S 6808mm⊗	10.0	40.1	(2.1	6.8	0.0	0.0	0.0	0.0	27.9	28.5	69.2) 	20.8	17.1				-%S 6815mm⊗	1	27.1	12.2
		Variety		8808-# = RZM-	8808 -10	-11	,	-12	-13	-14	-15	-16	-17	-18	-19	Checks 8833	8869		8848M	8810M	M6988	R836	US H11	8815-# = RZM-\$S	1	- 2	m I

EVALUATION OF MONOGERM S1 PROGENY LINES FROM RANDOM-MATED POPULATIONS, SALINAS, CA., 1999

(TESTS 899, 1899, 4899) (cont.)

Yield Sucrose RJAP 100' Score 1bs \frac{\partial}{\partial} \text{ Mean} \text{Mean} \text{ Mean} \	Sugar
	Υı
	ᆌ

(TESTS 899, 1899, 4899)

(cont.)

	Root	اید	١		4.	9.	4.	7.	ო.	۳.	4.	7 **
	8	Rot	∞		44	68.6	58.4	40.2	22.3	30.3	84.4	3.7** 1.7**
Rzm)	RZM	Resist	%		53.8	28.6	66.7	78.3	75.3	26.5	21.9	
Test 4899 (Rzm)		RJAP	%		83.2	83.9	86.5	75.3	82.0	3.3	2.5	2.5**
Test		Sucrose	₩		15.87	15.27	16.60	14.40	5455.6 15.57	1.07	4.25	1.5**3.69
	Sugar	Yield	1bs		4970	5493	3598	3896	5455.6	2209.2	25.2	1.5
	λλ	Score	Mean						5. و.	8.0	8.1	2.0**
(/	1001	No.						139.0	23.6	10.5	3.6**
Test 1899 (VY		RJAP	% [82.4	3.3	2.5	2.5**
Test 1		Sucrose	o⁄o [15.68	0.71	2.78	9.0** 7.25**
	ugar	Yield	1bs						5182.2	1279.9	15.2	*0.6
NB)		DM	%]		17.6	0.0	2.0	35.7	11.6	19.7	105.7	2.7**
Test 899 (NB)	Stand	Count	Mean	⊗	15.3	17.0	16.3	16.0	15.0	2.1	9.8	3.8**
Tes		%Bolt	8/26	%S 6821mm	30.7	3.9	69.4	13.9	24.2	17.6	45.1	13.9**
		Variety		8821-# = RZM-%S 6821mm⊗	8821 - 1	- 2	۳ ۱	- 4	Mean	LSD (.05)	C.V. (%)	F value

See Tests 1899 and 4899 for performance under virus yellows and rhizomania. TEST 899 NOTES:

Inoculated See test 899 and 4899 for companion tests under bolting and rhizomania conditions. TEST 1899 NOTES: See te with VY (BYV-BWYV-BChV).

Harvested by hand with 5% tare. Lifted roots counted (harvest count), rhizomania susceptible roots counted and % RZM resistant calculated, and root rot counted (Sclerotium rolfsii). All roots in plot weighed but only nonrotted roots used in sugar sample. Also see results from bolting, virus yellows, and TEST 4899 NOTES: Brawley trials.

HYBRID PERFORMANCE OF MONOGERM S1 PROGENY LINES, SALINAS, CA., 1999

(TESTS 999, 2099, 2399, 5599, B799)

B799(IV)	 %	13.08 11.60 11.08	12.32 11.81 10.52 11.68	12.63 11.86 11.68 11.99	12.49 11.24 11.89 11.74	11.22 11.76 11.85 12.31
Test B79 Sugar Yield S	١,	8420 7967 6872	8130 6185 6453 8029	7162 7193 6564 7298	6340 6650 7297 6658	6625 7972 7673 7742
m) RJAP	o%	85.9 87.0 84.4 86.0	87.0 85.5 86.6 86.5	85.4 86.6 85.4	86.2 84.5 83.7	86.1 85.6 86.7 87.5
5599 (Rzm) Sucrose F	o%	18.02 17.30 15.70 16.88	17.13 17.73 16.58 17.53	17.50 17.02 17.13	16.67 17.25 17.28	16.55 16.98 15.38
Test Sugar Yield	lbs	8093 9847 6663 6634	7963 9227 6091 8298	7001 6528 8374	6977 6424 7429	6707 7307 6853 5366
Sucrose	o%	16.55 16.84 16.04 15.79	16.05	16.41	16.08	15.51
Test 200 Sugar Yield	lbs	9866 11277 11154 10410	11727	11237	9184	0686
999 (NB)	%	2.2 6.0 22.8	34.2 10.3	10.2 15.9 2.1 9.0	0.0 3.7 13.2 23.4	19.7 21.1 8.8 3.9
Test 999 Bolting	% I	55.9 54.5 34.6	23.2 53.8 71.3	32.6 32.6 64.0 61.7	14.7 18.8 56.8 29.0	62.0 65.2 18.9 21.6
ld) RJAP	∞ I	84.4 85.2 84.8 83.9	83.2 82.4 83.1 83.1	83.9 83.0 83.6 84.5	83.2 83.5 82.9 81.8	84.0 84.7 83.8 84.7
2399 (Yield) Sucrose Ru	o%	17.44 17.98 16.49 16.81	16.50 17.25 16.19	16.69 16.16 15.94 16.25	16.25 16.49 16.65	15.76 16.09 15.48 16.01
Sugar Yield S	1	13365 15419 14788 14548	from popn-833 13681 13336 1 11503	14527 14224 12707 11976	from popn-834 12541 12710 2 12789 3 13397	13489 3 13107 from popn-828 9 13242 10 13665
Varietv		Checks Rifle B4776R Y869H50 Y869H46	S ₁ lines from Y869H35 Y869H5 Y869H33-1	Үвбэн33-10 Үвбэн33-11 Үвбэн33-12 Үвбэн12	S ₁ lines from Y869H29 Y869H34-1 Y869H34-2 Y869H34-3	Y869H34-5 Y869H34-8 S ₁ lines from Y869H28-9 Y869H28-10

(TESTS 999, 2099, 2399, 5599, B799)

(cont.)

B799 (IV)	Sucrose	o,∞		11.32	1.3	8	2.1	11.90	2.3	ا ا	11.55	11 57	. –		•	Ť.		2.3	1.8	11.91	2.0		2.3	1.8	12.74	2.6	1.4
Test B79				6469	8299	6926	7032	7300	4	(*)	7348	7569	0 6	6890	7 (4		10	27	7789	31		4	S	6750	0	σ
m)	RJAP	»۱		5	9	85.3	ъ.	8	4	g	85.2		. 4	85.2) (5	ر ا	87.1	9		ف	•	83.8		4.
5599 (Rzm)	Sucrose	o-01		6.9	7.0	17.22	6.7	ω.	6.3	7.3	16.92	7		. 6	: 5	r •		6.1	6.8	16.97	7.2	1	/ . 1	6.7	16.67	6.9	6.0
Test	Sugar Yield 8			53	82	7510	73	8037	4	87	8112	6647	75	N)	5777	r r		7895	80	81	88	1	1158	7975	7511	0669	8317
2099 (VY)	Sucrose	o∤≎l		15.79		15.20	5.9							5.6	16 10	•			15.64								
Test 20	Sugar	1bs		10644		10268	0							9676	10500				9585								
999 (NB)	ΜΩ	%		٠	ö	4.0	•	ω.	•	•	2.2	~	ď	14.1	ָ ע)			•	16.0	•	(٠	4	2.1	•	•
Test 99	Bolting	æI		о О	Э.	59.7	7.	و	9	17.1		· σ	0		. 4	•			4.	49.2	9			رى	45.2	4.	Ŋ.
14)	RJAP	o 0		84.3	85.8	84.0	82.3	84.3	83.4	83.0	84.2	85.9	84.0	79.7	84 6	;		83.5	•	•	83.8		83.2	82.8	•	•	83.7
2399 (Yield	Sucrose	o 0		•	16.21	Ŋ.	16.71	15.76	16.44	16.81	6.4	16.00	16.56	16.14	16.61	1 5 -		15.71	16.21	16.33	16.45	(ຖ	\sim	16.25	\leftarrow	0
ابد	Sugar Yield 8	1bs	698-udod	13219	13448	13923	12530	12721	12834	13278	14662	13928	13688	13485	12780)) !	from popn-836	12862	12829	13477	12558	월	13222	13012	13599	13713	12143
	Variety		S ₁ lines from	х869н69	х 869H69- 1	1	¥869H69- 4	х869н69- 5	9 -69н698Х	7 -69H698Y	х 869H69-13	X869H69-19	Y869H69-20	X869H69-20B	Y869H69-24		m	69H38	хв69н36- 3	Y869H36-11	X869H36-14	S ₁ lines from	T-//UEDOT	X869H77-1B	X869H77-2	X869H77-3	X869H77-4

HYBRID PERFORMANCE OF MONOGERM S1 PROGENY LINES, SALINAS, CA., 1999

(TESTS 999, 2099, 2399, 5599, B799)

(cont.)

B799 (IV)	Sucrose	%	I	1.3	11.67	0.5	1.5	1.6	1.9	11.97	1.8		ο.	11.29	1.1	1.1	11.90	1.3	1.3		2.2	12.23	2.3	3.1	1.3	10.78	2.7
Test B	Sugar	lbs		9689	7561	6736	9029	93	96	6401	49		36	7200	50	9	8824	7	_		73	7298	20	14	29	6337	41
2m)	RJAP	%	I	ω.	86.9	9	ъ.	7.	9	86.1	9			84.8	7.	4.	86.7	2	ъ.		ė.	87.6	4.	7.	5	87.2	9
5599 (Rzm)	Sucrose	l l	I	9.9	16.30	6.9	6.4	7.0	6.9	16.58	6.0			17.42	7.1	7.4	17.20	7.8	7.7		7.0	17.00	9.9	7.2	9.9	16.17	6.3
Test	Sugar Yield	lbs		63	7211	77	4	11	15	6812	91			8481	13	87	9505	79	57		σ	7065	4	0	N	7123	œ
2099 (VY)	Sucrose	o⁄(>	I		15.54	Ŋ.								15.69		15.77			16.04								
Test 2	Sugar Yield	1bs			10419	9839								9617		11481	10438		11896								
999 (NB)	MO		i	•	2.1	•	0	•	•	14.6	÷.		4	14.7	÷.	•	23.3	•	•		•	14.3	0.0	•	•	0.0	•
Test 99	Bolting	0,0	1	ω.	50.5	9	.	4.	2	26.3			Η.	22.9	÷.	•	23.2	•	œ œ		ъ.	61.7	<u>ი</u>	;	0	14.8	7.
1d)	RJAP	%	I	83.3	83.6	84.6	83.4	•		84.9	•		83.6	82.9	•	•	83.4	•	•		•	83.6	•	84.1	83.0	84.4	•
2399 (Yield)	Sucrose	%	I	9	15.41	9	9	9	9	16.40	വ	-4	6.3	16.38	6.1	6.3	16.19	6.5	6.2		16.14	16.67	16.81	16.09	5.8	15.90	6.4
الد	Sugar	1bs	Ď	12545	12537	13459	11858	13819	13888	13242	13886	popn-831	13614	13113	13739	14040	13328	12640	13953	popn-808	13826	13911	13697	12914	13890	13002	12542
	Varietv		S. lines from			X869H79-3		X869H79-5	X869H79-5B	X869H79-6	Y869H79-10	S ₁ lines from	¥869H4	Y869H27-1	Y869H27-2	Y869H27-7	X869H27-8	X869H27-9	X869H27-10	S ₁ lines from	Y869H9-1	х869н9-2	х869н9-3	X869H9-4		8-6Н698Х	

(TESTS 999, 2099, 2399, 5599, B799)

(cont.)

	Test	Test 2399 (Yield)	1d)	Test 999	999 (NB)	Test 2099 (VY)	(XA) 660	Test	. 5599 (Rzm	<u>н</u>	Test B799(IV)	(AI) 66,
	Sugar Yield	Sucrose	RJAP	Bolting	ΣΩ	Sugar	Sucrose	Sugar	Sucrose	R.TAD	Sugar	
	1bs	∞	∞	∞ 1	o-∞1	1bs	æ1	1bs	o~ 1	æ1	1bs	* Cons
FO H	lines from popn-808	\sim										
	12633	15.21	83.1	45.4	13.1			7438	15.73	86.5	8216	12 07
	13270	16.02	83.4	30.6	9.6			8208	17.33	· 1	6974	11 77
	12132	15.76	85.1	26.2	4.4			7038	15.65	86.5	5174	
rom	S ₁ lines from popn-818											
	13119	16.66	83.0	37.5	4.8			7552	17 00	ر م	9006	5
X869H15-2B	13648	16.21	82.3	20.4	5.9			8407	16.95	. 7 . 7	7907	11.01
									,	•	"	07.11
	13361	16.41	83.0	30.1	18.5			7803	16.85	1 78	6631	10 65
	12745	15.96	82.4	18.8	32.1			7794	16.80		10000	12.03
	13384	15.96	80.3	4.2	18.6			7777	16.00		0 4 0	•
X869H15-21	12687	15 07	7 00					r - r -		04.5	819/	12.44
	1	ñ .	7 0	y. V	9.17			8174	16.88	85.1	6928	12.48
	13279.1	16.29	83.3	36.0	12.1	10493.7	15.84	7 2177	28.0	α u	7102 2	70
	1899.2	0.71	2.3	22.7	19.1	927 0		1407) () ()	7.001.	99.11
	10.3	3.13	2.0		7 7					0.6	1342.3	
	1.1	.1NS2.89**		*	1 2NS	9 4	· · · · *	1.01	0 . k	7 . 7	13.4	5.79
		1	ſ	·)	C1177 - T	, 1	`	× × ○ · †	× 4.56××	1.5*	2.1	2.1**2.68**

SUGAR BEET RESEARCH

1999 REPORT

Section B

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Colorado Agricultural Experiment Station

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USDA-ARS-NPA Sugar Beet Research Unit's Mission Statement

Utilize distinctive site environmental and disease-free characteristics and specifically developed team expertise to: develop new knowledge and adapt biotechnologies to modify host-pathogen relations that affect disease resistance, pathogenesis, and epidemiology in sugar beet and other plant species pertinent to sugar beet cultivation; discover new information and techniques to identify and produce genotypes exhibiting superior disease and stress tolerance and agronomic qualities; and provide new knowledge that improves production efficiency and biochemical processing characteristics of sugar beet.

USDA-ARS -NPA COLORADO-WYOMING RESEARCH COUNCIL

The Sugar Beet Research Unit is a part of the Colorado-Wyoming (CO-WY)Research Council. This Council was chartered to promote and coordinate cooperative research activities among CO-WY Council research units; and facilitate communication and interaction with the Northern Plains Director, and among research programs and units and with customers locally, regionally, nationally and internationally. The five research units listed below publish an annual compilation of research reports. Many of the units are considering or have placed these reports on individual home pages which can be accessed through the NPA home page at www.npa.ars.usda.gov.

Rangeland Resource Research Unit (RRRU) - Cheyenne, WY, Fort Collins, CO & Nunn, CO MISSION STATEMENT: The mission of the Rangelands Resources Research Unit is to develop an understanding of the interrelationships of the basic resources that comprise rangeland ecosystems. Research is directed toward the development of science and technology that contributes to enhanced forage and livestock production and sustainable, productive rangelands in the Central Great Plains.

Central Plains Resources Management Research Unit (CPRMRU)- Akron CO.

MISSION STATEMENT: To enhance the economic and environmental well-being of agriculture by development of integrated cropping systems and technologies for maximum utilization of soil and water resources. Emphasis is on efficient use of plant nutrients, pesticides, and water and soil conservation/preservation.

Great Plains Systems Research Unit (GPSRU) - Fort Collins, CO.

MISSION STATEMENT: Help develop and implement sustainable and adaptive agricultural systems by: (1) synthesizing, quantifying, evaluating, and enhancing knowledge of processes; (2) developing integrated models of agricultural systems; (3) providing technology packages to agricultural communities and action agencies.

Soil-Plant-Nutrient Research Unit (SPNRU) - Fort Collins, CO.

MISSION STATEMENT: To develop and evaluate new knowledge required to efficiently manage soil, fertilizer and plant nutrients (emphasis on nitrogen) to achieve optimum crop yields, maximize farm profitability, maintain environmental quality and sustain long-term productivity.

Water Management Resources Unit (WMRU) - Fort Collins, CO.

MISSION STATEMENT: Research emphasis is to integrate applied and basic principles to develop improved water, chemical, and alternative weed management systems and irrigation system designs. Improvements are directed toward sustainable, environmentally sound and efficient systems based on soil, water, fertility, energy, and weed ecology principles. This encompasses understanding physical and biological phenomena and developing computer simulation models and precision farming systems to transfer new technologies to producers, consultants, action agencies, industry, and scientists.

For a copy of the Colorado-Wyoming (CO-WY)Research Council Annual Report or information on any of these programs, please note the following contacts:

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PUBLICATIONS & ABSTRACTS

- 1. Panella, L. Long Term Performance of Artificially Inoculated Cercospora Leaf Spot Nurseries. pp. 123, *In*: xyz (eds.) Cercospora. Advances in Sugar Beet Research, vol.2, IIRB, Brussels, Belgium. 1999. (in press)
- 2. Panella, L. and L. Frese. Cercospora resistance in *Beta* species and the development of resistant sugar beet lines. pp. 123, *In*: xyz (eds.) Cercospora. Advances in Sugar Beet Research, vol.2, IIRB, Brussels, Belgium. 1999 (in press)
- 3. Panella, L. Evaluation of *Beta* PIs from the USDA-ARS NPGS for resistance to Cercospora leaf spot, 1999. Biological and Cultural Test for Control of Plant Diseases. Am. Phytopathol. Soc. Vol.15: 2000. (Accepted 12/10/99).
- 4. Panella, L. Evaluation of *Beta* PIs from the USDA-ARS NPGS for resistance to curly top virus, 1999. Biological and Cultural Test for Control of Plant Diseases. Am. Phytopathol. Soc. Vol.15: 2000. (Accepted 12/10/99).
- 5. Panella, L. Evaluation of *Beta* PIs from the USDA-ARS NPGS for resistance to Rhizoctonia root rot, 1999. Biological and Cultural Test for Control of Plant Diseases. Am. Phytopathol. Soc. Vol.15: 2000. (Accepted 12/10/99).
- 6. Panella, L. Evaluation of Rhizoctonia-root-rot-resistant germplasm released by the USDA-ARS Sugar Beet Research Unit over 30 years, 1999. Biological and Cultural Test for Control of Plant Diseases. Am. Phytopathol. Soc. Vol.15: 2000. (Accepted 12/10/99)
- 7. Panella, L. USDA-ARS Sugar Beet Research at Fort Collins In it for the Long Haul! Sugar J. 11: March, 2000. (Popular Press)
- 8. Panella, L. Screening Sugar Beet Germplasm for Rhizoctonia Root Rot Resistance. Agr. Abstr. p. (ASA-CSSA-SSSA Annual Meeting, 31 Oct 4 Nov, Salt Lake City, UT). 1999. (poster)
- 9. Wickliffe, E., Otto, K., Schwartz, H.F., Brick, M. A., Ogg, B., Byrne, P., Fall, A., Panella, L., and Hill, A. Fusarium will variability in dry bean and sugarbeet. 15th Biennial Meeting of the Bean Improvement Cooperative, Calgary, Canada, Nov. 8-12, 1999. (poster)
- 10. Weiland, John J., Robert T. Lewellen, J. Mitch McGrath, Lee Panella, and Ming H. Yu. Tagging of disease resistance genes in sugarbeet (*Beta vulgaris* L.) with molecular genetic markers. Plant & Animal Genome VIII Conference, San Diego, CA, January 9-12, 2000. (Abstract)

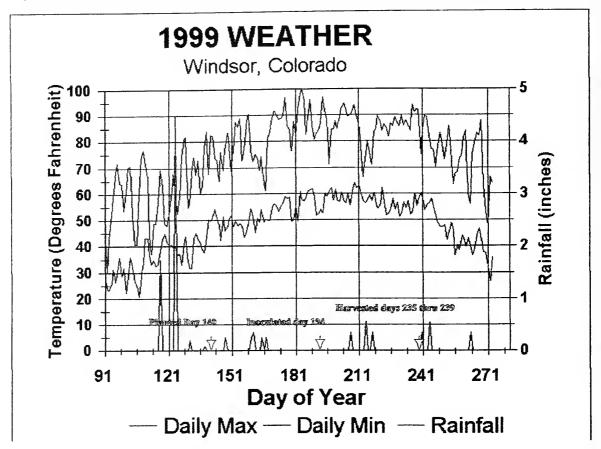
EVALUATION OF CONTRIBUTED LINES FOR RESISTANCE TO RHIZOCTONIA SOLANI, A CAUSAL FUNGUS OF SUGAR BEET ROOT ROT. (BSDF Project 903) L. Panella

Annually, for over thirty years, the breeding program in Fort Collins has created an artificial epiphytotic through inoculation with *Rhizoctonia solani* to evaluate and select for resistance to root rot caused by this pathogen. We have been pleased to participate and lead this cooperative research project between the ARS, Colorado State University, and the BSDF. The project primarily involved field studies conducted on 35 acres of leased land near Windsor, CO. Randomized, complete-block designs with five replicates were used to evaluate Fort Collins ARS breeding germplasm. Rhizoctonia-resistant line FC703 and highly susceptible FC901/C817 were included as internal controls, along with highly resistant FC705-1.

One-row plots, planted May 20th, were 12 feet long with 22 inches between rows and 8-10 inches within-row spacing. Inoculation with dry, ground, barley-grain inoculum of *Rhizoctonia solani* isolate R-9 was performed on July 13th; immediately after inoculation, a cultivation was performed so as to throw soil into the beet crowns. The field was sprayed twice with Betamix Progress, Upbeet, and Stinger (June 2 and 12) to control weeds. The field was thinned by hand and irrigated as necessary. Beets were harvested August 23 through 27. Each root was rated for rot on a scale of 0 to 7 (dead) as previously described. ANOVAs were performed on disease indices (DIs), percent healthy roots (classes 0 and 1 combined), and percentage of roots in classes 0 thru 3. Percentages were transformed to arcsin-square roots to normalize the data for analyses. LSDs are provided for comparing entries with those of our internal checks.

We also had just a little rain in the week after planting with warming temperatures (Figure 1). Therefore, stands were excellent and the 1999 Rhizoctonia epidemic started strong and progressed quickly, becoming severe by mid August. Differences in DIs among entries in all tests were highly significant (P < 0.001). Mean DIs across all tests for highly resistant FC705-1, resistant FC703, and the highly susceptible check were 3.3, 3.9, and 6.2, respectively. Percentages of healthy roots were 17.8, 9.5, and 0.5 for these internal controls. Percentages of roots in disease classes 0 thru 3 were 56.3, 38.0, and 4.0, respectively. The highest and lowest DIs for evaluated lines were 6.8 and 2.0, respectively.

USDA-ARS 1999 Rhizoctonia Disease Nursery, Fort Collins, CO.



<u>Figure 1 & Table 1.</u> 1999 Rhizoctonia Root Rot Nursery, Fort Collins, CO. The Graph above summarizes the weather data for our Rhizoctonia Root Rot Nursery in 1999. The table below presents summary data of the entire nursery. The experiment mean, the mean of the susceptible check, the mean of the resistant check, and the mean of the highly resistant check are given for each of the experiments in the nursery. LSD is at the t=0.05 level.

		Dis	ease Ir	idex		Perce	nt Hea	lthy (c	lasses (8 1)	Pe	rcent i	n Clas	ses 0 to	3
Exp.	Mean	Sus.	Res.	H Res.	LSD	Mean	Sus.	Res.	H Res.	ESD	Mean	Sus.	Res.	H Res.	SI
1R	5.0	6.2	3.8	3.4	80	8	0	8	22	13.2	21	6	44	48	
3R	4.6	6.0	3.6	3.5	75	5	0	6	16	11,5	22	2	50	58	
4R	4.1	5.9	3.8	3.3	90	13	0	12	22	14.5	39	2	42	56	
5R	5.7	6.4	4.2	3.1	.63	3	0	8	28	9.4	9	4	30	58	
7R	5.2	6.5	3.9	2.9	.69	5	0	16	26	10.6	13	2	34	64	
8R	5.6	6.2	4.1	3.4	67	2	2	10	0	8.5	9	8	32	56	
9R	4.7	6.2	4.0	3.8	.73	9	0	4	0	9.1	28	4	30	40	15
10R	4.2	6.0	3.7	2.8	.87	12	2	12	28	14.9	36	4	42	70	18
Avg.	4.9	6.2	3.9	3.3		7.1	0.5	9.5	17.8		22.1	4.0	38.0	56.3	

Percent in Classes is the transformed value (arcsin-square root)

Mean = Experiment Mean;

Sus. = Susceptible Check (FC901/C817);

Res. = Resistant Check (FC703);

H Res. = Highly Resistant Check (FC705/1)

EVALUATION OF CONTRIBUTED LINES FOR RESISTANCE TO CERCOSPORA BETICOLA, CAUSAL FUNGUS OF CERCOSPORA LEAF SPOT (BSDF Project 904) L. Panella

The breeding program in Fort Collins has created an artificial epiphytotic through inoculation with *Cercospora beticola* annually for over forty years to evaluate and select for resistance to leaf spot caused by this pathogen. We have been pleased to participate and lead this cooperative research project between the ARS, Colorado State University, and the BSDF. The project primarily involved field studies conducted on 35 acres of leased land near Windsor, CO.

Randomized complete-block designs, with three replications were used to evaluate commercial and experimental entries. Internal controls included a highly susceptible synthetic and a resistant check (FC 504/502-2//SP6322-0). The nursery was planted on April 29th. Fertilization was 75% of the soil test recommendation to minimize leaf growth, which can interfere with visual evaluations. Differences among lines were highly significant in all tests at each of three evaluation dates. Two-row plots were 12 feet long, with 22-inch row spacing and an 8 - to 10-inch within-row plant spacing. The trial was planted on April 20th in Windsor, CO. Inoculation was performed on June 30th and again on July 7th. Evaluations were made on September 7th, 14th, and 22nd, with the peak of the epidemic occurring on or about the last date. The field was sprayed twice with Betamix Progress, Upbeet, and Stinger (May 14th and 24th) to control weeds. The field was thinned by hand and irrigated as necessary.

We had good spring rain in 1999 and emergence was excellent and we got off to an early start. The 1999 leaf spot epidemic started strong and progressed rather slowly, but eventually became more severe by late August. We had a period between of about one month right after inoculation, in which we had relatively high evening temperatures (Figure 2), which helped disease development, however by September or evening temperatures had dropped. At our third evaluation, means of the resistant and susceptible internal controls were 3.1 and 6.4 (scale of 0-10), respectively, across the nursery. In 1998 (September 8), these means were 3.2 and 5.3, respectively. Means of contributor lines on September 22 ranged from 2.7 to 9.0, compared with 2.5 to 8.0 in the milder epidemic of 1998.

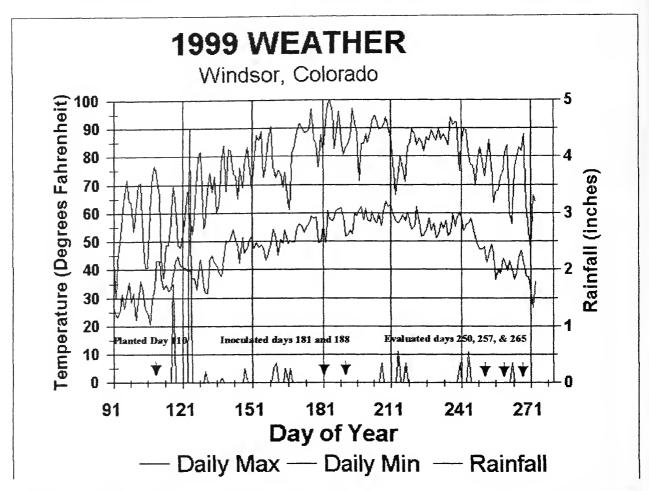


Figure 2 & Table 2. 1999 Cercospora Leaf Spot Nursery, Fort Collins, CO. The Graph above summarizes the 1999 weather data for our Cercospora Leaf Spot Nursery int 1999. The table below presents summary data of the entire nursery. The experiment mean, the mean of the susceptible check, and the mean of the resistant check are given for each of the experiments in the nursery, for each evaluation date. The highest mean rating given on September 22nd was a 9.00 and the lowest a 2.67.

		Septen Disease	nber 7 th e Index			Septem Diseas	ber 14 ⁶ e Index			Septem Diseas	ber 22° e Index	
Exp.	Mean	Sus.1	Res. ²	LSD	Mean	Sus.	Res.	LSD	Mean	Sus.	Res.	LSD
1A	4.2	5.0	2.8	0.97	4.7	5.2	6.5	1.08	5.3	6.5	2.8	31.1
2A ³	4.5	4.5	2.3	1.97	5.3	6.5	2.5	1.75	5.9	7.0	3.5	1.5
3A	4.4	4.8	2.5	1.24	5.1	5.3	3.5	0.99	5.6	5.5	3.5	0.9
4A	4.7	5.8	2.7	0.81	5.3	6.3	2.5	1.10	5.5	6.7	2.8	0.8
5A	5.0	5.2	3.3	0.93	5.4	5.8	3.2	0.80	6.0	6.2	3.7	0.6
6A	4.1	4.8	2.7	0.77	4.8	5.7	3.0	0.90	5.3	6.3	3.2	0.9
7A	3.7	5.8	2.8	0.77	3.9	6.0	2.8	0.93	4.5	6.7	3.3	10
7A⁴	3.6	5.0	2.7	0.77	3.9	5.2	2.8	0.93	4.7	6.5	3.3	10
8A	3.6	5.2	2.8	0.83	3.6	5.7	2.5	0.92	4.0	6.3	2.7	1.1
9A	3.2	5.0	2.3	0.77	3.3	5.7	2.0	0.92	3.9	6.3	2.7	1.0
Mean	4.10	5.11	2.70	6.5	4.53	5.73	3.13		5.07	6.41	3.14	1000

¹Cercospora Susceptible Check - SP351069-0

²Cercospora Resistant Check - FC 504CMS/FC 502-2//SP6322-0

³There were only two replications of Experiment 2A

⁴There were two separate tests in Experiment 7A

RHIZOCTONIA ROOT ROT RESISTANCE AND DEVELOPMENT OF GENETIC RESISTANCE IN SUGAR BEET - BSDF Project 440

L. Panella

This facet of the USDA-ARS Fort Collin's sugar beet breeding program has as its goals: 1) the understanding the genetics of the *Rhizoctonia solani*/sugar beet interaction in order to better facilitate development of germplasm with high levels of resistance to Rhizoctonia and other sugar beet diseases, and 2) to provide the knowledge to better manage this disease in sugar beet production areas. It is an integrated research program with greenhouse, laboratory, and field components. Genetic information developed previously in our research is used to execute additional cycles of pathogen inoculation, plant selection, and recombination among germplasms that we have in our cyclic improvement program. Germplasms in various stages of improvement are evaluated for resistance in inoculated field tests. Results of these tests form the basis of decisions about specific germplasm, i.e., retain, shelve, discard, recombine, release, etc. Germplasms likely to be useful for variety improvement are identified and released for use by other sugar beet breeders.

1999 Field Research on Rhizoctonia Root Rot of Sugar Beet.

The breeding program in Fort Collins has created annually an artificial epiphytotic through inoculation with *Rhizoctonia solani* for over thirty years to evaluate and select for resistance to root rot caused by this pathogen. We have been pleased to participate and lead this cooperative research project between the ARS, Colorado State University, and the BSDF. The project primarily involved field studies conducted on 35 acres of leased land near Windsor, CO. Randomized, complete-block designs with five replicates were used to evaluate Fort Collins ARS breeding germplasm. Rhizoctonia-resistant line FC703 and a highly susceptible check (FC901/C817) were included as internal controls, along with highly resistant FC705-1.

One-row plots, planted May 20th, were 14 feet long with 22 inches between rows and 8-10 inches within-row spacing. Inoculation with dry, ground, barley-grain inoculum of *Rhizoctonia solani* isolate R-9 was performed on July 13th; immediately after inoculation, a cultivation was performed so as to throw soil into the beet crowns. The field was sprayed twice with Betamix Progress, Upbeet, and Stinger (June 2 and 12) to control weeds. The field was thinned by hand and irrigated as necessary. Beets were harvested August 23 through 27. Each root was rated for rot on a scale of 0 to 7 (dead) as previously described. ANOVAs were performed on disease indices (DIs), percent healthy roots (classes 0 and 1 combined), and percentage of roots in classes 0 thru 3. Percentages were transformed to arcsin-square roots to normalize the data for analyses ("Z% Hlthy" and "Z% 0-3" in the accompanying tables). Both percentages and arcsins are given in the table, and LSDs are provided for comparing arcsins of your entries with those of our internal checks.

We also had just a little rain in the week after planting with warming temperatures. Therefore, stands were excellent and the 1999 Rhizoctonia epidemic started strong and progressed quickly, becoming severe by mid August. Differences in DIs among entries in all tests were highly significant (P < 0.001). Mean DIs across all tests for highly resistant FC705-1, resistant FC703, and the highly susceptible check were 3.3, 3.9, and 6.2, respectively. Percentages of healthy roots were 17.8, 9.5, and 0.5 for these internal controls. Percentages of roots in disease classes 0 thru 3 were 56.3, 38.0, and 4.0, respectively. The highest and lowest DIs for evaluated lines were 6.8 and 2.0, respectively.

Allotment of Fort Collins "FC" numbers (3-digit numbers)

"FC" numbers are "convenience" numbers for "seed releases" or purposes where a permanent line designation is needed — i.e. a number that does not change from generation to generation where little or no selection pressure is applied. Initially, an "FC" no. was written thus "FC 501" [now FC727], "FC 502 CMS" [now FC715CMS], etc. Sublines (from selfing) were designated thus, "FC 502/2" [now FC709-2], "FC502/3" [now FC502-3], etc. The same applies when the line is substantially changed by selection without selfing.

Below 500	Originally LeRoy Powers - now parental lines and special genetic stocks.
500's	Leaf Spot Resistant (LSR), Type-O lines & male steriles [CMS]
600's	LSR-Curly Top Resistant (CTR), type-O lines & male steriles [CMS]
700's	Rhizoctonia Resistant
800's	LSR-CTR-Rhizoctonia resistant
900's	Pollinators, LSR-CTR type

This year, I also completed a one year evaluation of most of the Rhizoctonia-resistant lines released from the USDA-ARS breeding project at Fort Collins (Table 3). This is a test from 1999 under the same conditions as the other contributor lines in this year's test.

Transforming Rhizoctonia-Resistant Populations to Germplasm with Multiple Disease Resistance

Root rot and leaf spot are two serious diseases of sugar beets caused by fungi (*Rhizoctonia solani* and *Cercospora beticola*, respectively). The diseases caused by these fungi may produce a severe reduction of yield in many sugar beet production areas. Cultural control measures are not adequate by themselves, and often no chemicals are registered for control of these diseases, or chemical control is expensive or environmentally unsafe. Increased levels of genetic resistance in sugar beet varieties are needed to minimize growers' losses from these diseases. In a hybrid crop like sugar beets, it is preferable that all of the parents contain some level of resistance to diseases prevalent in the area in which the hybrid is to be grown. Multiple disease resistance is a difficult goal in a crop improvement program, especially when working with an outcrossing species. In alternating generations of selection, some of the progress made in resistance to one disease is lost while selecting for resistance to other diseases.

One way of solving the problem of selecting for multiple disease resistance is the use of progeny testing. By testing the progeny of individual mother roots, plants with multiple disease resistance can be identified and used as parents of the next generation. The most efficient use of progeny testing is when the genotype of both parents is controlled, and the most effective way to do this is through self-pollination. In sugar beet, there is a dominant, self-fertility gene that permits self-

pollination. Used in conjunction with genetic male sterility, to insure cross pollination, a system of selfed-family progeny testing can be utilized.

This effort is based on the Rhizoctonia-resistant materials from the programs of John Gaskill and Richard Hecker, and disease resistant germplasm from other sources to produce germplasm highly resistant to Rhizoctonia solani. This base of Rhizoctonia-resistant germplasm is being combined with material from the USDA-ARS breeding programs at Salinas and Fargo, as well as with sources for higher yield and sucrose. The Salinas material has the self-fertility allele, is segregating for genetic male sterility, and also contains a broad spectrum of resistance to diseases of importance in California as well as other sugar beet production areas (including rhizomania, powdery mildew, virus yellows, and curly top virus). Fargo sources of root maggot and Cercospora leaf spot resistance are also being utilized.

A number of source populations are being developed. The germplasm, FC712(4X) has been released in 2000. This germplasm was developed in our research project that has been contributed to, in kind, by the Beet Sugar Development Foundation. This tetraploid pollinator germplasm combines excellent Rhizoctonia-root-rot resistance with a good level of Cercospora leaf spot resistance. Populations whose development was begun under the breeding program of Dr. Richard Hecker are still being evaluated and selected in the field. These germplasms and other germplasms from the Fort Collins program were field-tested in summer of 1999 for resistance to *R. solani* (Tables 3-4), *C. beticola* (Tables 5-7), and the curly top virus (Table 8). More germplasms that were selected for increased resistance to Rhizoctonia-root-rot in 1998, and tested in 1999, will be tested again in 2000; and the most promising of these will be released in the future.

There currently are four major groups of Rhizoctonia-resistant germplasms currently under development.

- 1. Germplasms developed in Dr. Hecker's breeding program for resistance to Rhizoctonia root rot and Cercospora leaf spot are being field tested and selected in the Rhizoctonia root rot nursery at Fort Collins (also in the Cercospora leaf spot and curly top nurseries).
- 2. Rhizoctonia-resistant monogerm polycross base population developed by a cross between FC708 and two Salinas germplasms, 2890 and 2859.
 - A. 2890 (sp) 0790 mm aa x 1890 (Salinas); is seed from aa plants open pollinated by A- plants. 0790 = population-790 cycle 5 synthetic by S₁ progeny, M.S. mm, O-type, good combining ability, adapted to California, S^f,. 1890 = BC population to population 790 to get Rz equivalent, remains variable for M-:mm, Rz-:rzrz, etc.
 - B. 2859 m (sp) = 1859, 1859R aa x A- (Salinas); Released in 1992 as C859. Sf, similar to 2890, but should have higher curly top resistance (CTR). Segregates and variable for M-:mm, Rz-:rzrz, A-:aa, predominant background is lines like C563, which is widely used in western USA as source of CTR, mm, O-type.
- 3. Rhizoctonia root rot resistance multigerm base population developed by a cross between FC709-2 and a Salinas germplasms, 2915.
 - A. 2915 (sp) RZM 1915-#m 1913-# aa x A (Salinas); Seed harvested from aa (ms) plants open-pollinated by A- (fertile) plants. This population will segregate for A-:aa, Rz-:rzrz, s*s*:sf-,

(>½ s^f), R-:rr, It will be multigerm, have moderate to good tolerance to virus yellows, curly top, bolting, Erwinia; variable for reaction to powdery mildew, production traits. Individual plants will be either As or aa. Background of population is mostly from OP, MM lines such as C46, C37.

4. Combination Rhizoctonia root rot and Cercospora leaf spot resistant multigerm pollinator population from FC907 (out of Fargo) and FC709-2.

Progress in 1999

- 1. Selections have been made in these populations and they have been crossed with other germplasm in a continuing *Rhizoctonia*-resistance breeding effort. One tetraploid multigerm pollinator [FC712 4(X)] has been released. It has excellent resistance to Rhizoctonia root rot and good Cercospora resistance. Three to five monogerm O-type lines with and without and CMS equivalents, selected in the 1996 Rhizoctonia nursery were re-tested this year and will be considered for release this summer.
- 2. S₁ families selected for curly top resistance from this monogerm base populations were selected in the Rhizoctonia nursery last year. This germplasm has been harvested increased in the Greenhouse at Fort Collins. This seed was planted in the mother root nursery at Fort Collins for increase and a split sample was sent to Salinas where it is being selected to see if the Holly gene for Rhizomania resistance is still segregating in the population. Rhizomania resistant plants will be intercrossed and seed planted in the Rhizoctonia nursery to be selected for resistance.
- 3. Individual selfed & half-sib families were harvested and progeny tested in the Rhizoctonia and curly top nursery in 1998 and Rhizoctonia nursery this year. Selections were made from the Rhizoctonia nursery and remnant seed is available for the top performers in the curly top nursery. These selections have been recombined and will be tested next year and the following year.
- 4. Seed, increase from Rhizoctonia-resistant selected roots of FC907 ((FC701 x FC607)BC₄), was tested in the Rhizoctonia and Cercospora nurseries next year. Selections made in a (FC709-2 x FC907)F₂ population in the Rhizoctonia were increased in the greenhouse and tested in the Rhizoctonia and curly top nurseries. This population will be re-selected in the Rhizoctonia nursery and then tested in the Rhizoctonia, Cercospora, and curly top nurseries. Half-sib family selections from this population (35 families) were made in the 1999 Cercospora nursery.

Future laboratory research will use the information gained from studying the pathogen *Rhizoctonia solani* to begin to look at the sugar beet reaction to this pathogen. Biocontrol work will resume once a new Research Plant Pathologist is on board.

Table 3. Experiment 4R, 1999. Rhizoctonia Resistance Evaluation of USDA-ARS Breeding Lines from Fort Collins CO, (Lee Panella)

981009H 931017 991011 991011 991012HO 961012HO 961012HO 991002MS 991002PF EL 50 WC980439 EL 52 99A003 EL 48 FC701 FC701 FC701 FC701-6 FC701-6 FC702-6 811055H FC702-6 811055H FC702-6 811055H FC702-6 811055H FC702-6 811055H	The state of the s	Description	5	/0 I HELLY	200		0 0 0
4 3 9 7		,ds7	0.30			14.5	18.2
4 2 9 9 2	T	(907/709-2)F2-Sel Rhzc	5.0	œ	12	13	18
4 2 9 9 7		Susceptible Check - (FC901/C817)	5.9	0	7	0	4
4 5 9 5			3.0	5 8	64	30	53
4 5 9 5			4.0	16	32	20	34
4 5 9 5	우	FC712/Mono-Hy A4	4.4	ထ	30	우	32
4 2 9 9 7	ş	FC712/Mono-Hy A4	4.9	7	14	4	19
4 v o o = 2	NS		5.6	7	ω	4	15
4 2 9 9 7	노		5.7	7	ထ	4	13
4 2 9 9 7	439		4 .8	œ	22	13	27
4.6.0		98J26-052	5.0	4	ω	7	15
4 7 9 9 7			5.5	4	9	7	4
4 rv o o -			5.7	0	9	0	7
იი ი -	T		3.6	18	44	24	41
o o -			4.7	4	18	7	22
9 +	T		3.2	20	56	23	48
9 +	Ó		5.4	4	12	ĸ	13
, -	T		3.2	22	56	27	49
_	T	Resistant Check	ა. 8.	12	42	18	40
		Highly Resistant Check	3.3	22	56	27	49
	T		3.3	22	20	25	45
FC708 831085HO	우		4.4	9	30	7	30
	_		2.5	32	80	34	99
FC709-2 921024		Fort Collins release	2.0	42	92	40	77
			3.6	14	54	19	20
(4X)		FC710 colchicine doubled	3.4	10	99	12	61
			4.9	7	22	4	22
2	T	Fort Collins Release	5.6	28	9/	31	64
(4X)		FC 712 colchicine doubled	3.0	56	62	27	55
່ ເ	우		4.3	16	32	21	33
			3.2	18	64	22	54
FC717 911031			6.4	4	16	7	21
FC718 911032			3.8 9.	14	42	21	40
FC719 911037			3.1	18	99	22	58

Table 3. Experiment 4R, 1999. Rhizoctonia Resistance Evaluation of USDA-ARS Breeding Lines from Fort Collins CO, (Lee Panella)

10	Seed Source	Description	DI,	% Hithy²	% 0 - 33	Z% ⁴ Hithy	Z% 0 - 34
		FSD.	06.0		Ť	14.5	18.2
FC720-1	961015	C718/(C718/FC708)	4.1	ထ	38	9	38
FC722-1	961010HO	C718/FC708	0.4	မ	44	; =	41
FC722CMS	961010HO1	C718/FC708	4.6	2	4	. 4	. 1
FC723	951016HO	EL44/FC708 mm	ထ	16	40	- 88	- œ
FC723CMS	951016HO1	EL44/FC708 CMS	6	. c c	36	<u>ተ</u>	2 6
FC724-1	961014	FC702/LSR-CTR	3.1	16	62	<u>6</u>	. v.
FC725	921008		ა ფ	22	62	27	52
FC726	931010		3.5	16	50	. 5	44
FC727	951017	Fort Collins release	4.1	9	388	16	80
FC728	921025		3.0	18	99	25.	, r.
FC729	921019	FC712/A4, 3 cycles Rhizoc, MM	6	12	40	5 4	8 8
FC907-1	971020	FC607/FC701 BC4 - 1 cycle of RhzcR sel	6.1	! o	, 4) c	- -
		Experiment Mean	4.1	13	38	16	37
Disease Inde	x is based on a s	Disease Index is based on a scale of 0 (=healthy) to 7 (= plant dead)					
² Percent of he	salthy roots (dise	Percent of healthy roots (disease classes 0 and 1 combined).					
³ Percent of dis	seased roots like	³ Percent of diseased roots likely to be taken for processing (disease classes 0 through 3 combined).	ondh 3 c	ombined).			
⁴ Percentages	were transforme	Percentages were transformed to arcsin-square roots to normalize the data for analyzes.	nalyzes.				
°P=0.05							

Table 4. Experiment 10R, 1999. Rhizoctonia Resistance Evaluation of USDA-ARS Breeding Lines, Fort Collins, CO; East Lansing, MI; and Fargo, ND.

П		Г																				_						
L% U - 3.	18,9	5 6	=	23	20	19	14	27	19	23	24	13	22	တ	4	39	55	52	53	29	20	28	20	59	7	09	40	34
7%. Hituk	14.9	31	0	တ	4	ഹ	7	13	7	13	13	7	S.	0	0	_	_	27	59	32	24	37	23	30	4	25	15	14
% U - 3		89	ဖ	24	18	4	10	56	14	16	20	12	20	ဖ	7	36	62	62	64	84	58	20	58	72	4	20	42	36
70 milliy		28	0	9	7	4	4	œ	9	80	œ	9	4	0	0	ဖ	ဖ	22	24	28	20	36	20	26	2	28	12	12
JUI	0.87	3.0	5.8	5.0	6.4	5.2	4 .	4.5	4.7	4.7	4.4	5.4	4.6	0.9	6.2	4.1	3.6	3.1	2.8	2.5	3.1	2.6	3.3	2.8	6.0	2.8	3.7	4.2
Location		East Lansing	East Lansing	East Lansing	East Lansing	East Lansing	East Lansing			East Lansing	East Lansing	Fargo	Fargo	Fargo	Fargo	Fargo	Fort Collins	Fort Collins	Fort Collins	Fort Collins	Fort Collins	Fort Collins	Fort Collins	Fort Collins	FC901/C817	FC705/1	FC703	
Seed Source		WC980435L										97N0050	96N0021	96N0022	98N0058	96N0023	971017	971018	881032H	921024	951017	961014	961015	891033	931017	831083	751080H	
Description		EL 51	99302-00	99J19-00	99320-00	99J25-024	98J26-2	98J26-3	98J26-7	98J25-38-5	98J25-01-3	F1001	F1002	F1004	F1005	F1006	FC712(4X)	FC710(4X)	FC712	FC709-2	FC727	FC724	FC720	FC710	Susceptible Check	Highly Resistant Check	Resistant Check	Experiment Mean

¹Disease Index is based on a scale of 0 (=healthy) to 7 (= plant dead).

²Percent of healthy roots (disease classes 0 and 1 combined).

³Percent of diseased roots likely to be taken for processing (disease classes 0 through 3 combined).

⁴Percentages were transformed to arcsin-square roots to normalize the data for analyzes.

⁵P=0.05

CERCOSPORA LEAF SPOT RESEARCH AND BREEDING FOR CERCOSPORA AND CURLY TOP RESISTANCE - (BSDF Project 441)

L. Panella

This element of the breeding program at Fort Collins is devoted to the development of germplasm with resistance to more than one sugar beet disease and improved agronomic characteristics. It is built on germplasm developed at Fort Collins over the last fifty years for combined resistance to Cercospora leaf spot and the curly top virus. This is an integrated breeding program with greenhouse and laboratory studies, and a field program based on testing in an artificial epiphytotic created in the unique Fort Collins environment. It involves close collaboration with the other USDA-ARS sugar beet programs in the U.S. and sugar beet seed industry customers. The major goals of this program are: 1) the development of sugar beet germplasm with resistance to more than one disease and excellent agronomic characteristics; 2) the improvement of breeding techniques, traditional and molecular, to develop this germplasm; and 3) an increased understanding of the sugar beet/pathogen interactions to improve management practices of these diseases in sugar beet production areas. Genetic information developed during this research will be used to execute additional cycles of pathogen inoculation, plant selection, and recombination among germplasms that we have in our leaf spot improvement program. Results of these tests will be the basis of decisions about specific germplasm, i.e., retain, discard, recombine, release, etc. Germplasms likely to be useful for variety improvement will be identified and released for use by other sugar beet breeders.

Increased resistance to Cercospora continues to be an extremely important goal. If the level of resistance available in most Cercospora-resistant experimental lines were present in commercial hybrids (along with good sugar and seed yield), the need for fungicides would be greatly reduced. That continued improvement in genetic resistance to this serious pathogen is still needed is evident by the occurrence of Cercospora strains that are resistant or increasingly tolerant to our most potent fungicides. Additionally, some of these fungicides may be removed from the market because of their perceived or real threat to the environment. In many areas where Cercospora leaf spot is a problem, the curly top virus also causes significant losses. And, there are some growing areas in which combined resistance to Cercospora leaf spot, Rhizomania, curly top, Rhizoctonia root rot, and other diseases are desirable. Germplasm is needed with combined resistance to these diseases, along with good combining ability for yield components.

1999 Field Research on Cercospora Leaf Spot of Sugar Beet

The breeding program in Fort Collins has created an artificial epiphytotic through inoculation with *Cercospora beticola* annually for over forty years to evaluate and select for resistance to leaf spot caused by this pathogen. We have been pleased to participate and lead this cooperative research project between the ARS, Colorado State University, and the BSDF. The project primarily involved field studies conducted on 35 acres of leased land near Windsor, CO.

Randomized complete-block designs, with three replications were used to evaluate commercial and experimental entries. Internal controls included a highly susceptible synthetic and a resistant check (FC 504/502-2//SP6322-0). The nursery was planted on April 29th. Fertilization was 75% of the soil test recommendation to minimize leaf growth, which can interfere with visual evaluations. Differences among lines were highly significant in all tests at each of three evaluation

dates. Two-row plots were 12 feet long, with 22-inch row spacing and an 8 - to 10-inch within-row plant spacing. The trial was planted on April 20th in Windsor, CO. Inoculation was performed on June 30th and again on July 7th. Evaluations were made on September 7th, 14th, and 22nd, with the peak of the epidemic occurring on or about the last date. The field was sprayed twice with Betamix Progress, Upbeet, and Stinger (May 14th and 24th) to control weeds. The field was thinned by hand and irrigated as necessary.

We had good spring rain in 1999 and emergence was excellent and we got off to an early start. The 1999 leaf spot epidemic started strong and progressed rather slowly, but eventually became more severe by late August. We had a period between of about one month right after inoculation, in which we had relatively high evening temperatures (Figure 2), which helped disease development, however by September or evening temperatures had dropped. At our third evaluation, means of the resistant and susceptible internal controls were 3.1 and 6.4 (scale of 0-10), respectively, across the nursery. In 1998 (September 8), these means were 3.2 and 5.3, respectively. Means of contributor lines on September 22 ranged from 2.7 to 9.0, compared with 2.5 to 8.0 in the milder epidemic of 1998.

Cercospora/Curly Top-Resistant Populations with Resistance to Multiple Sugar Beet Diseases and Superior Agronomic Characteristics

Advanced breeding lines or Cercospora-resistant germplasms from Fargo (5), Salinas (16), East Lansing (10), and Fort Collins (9) were evaluated in Experiment 7A at the ARS leaf spot nursery at Ft. Collins (Table 5). A blend of resistant and susceptible commercial hybrids was also evaluated by Larry Campbell - USDA-ARS at Fargo, ND. An additional 26 Fort Collins advanced breeding lines or released germplasms were evaluated for Cercospora leaf spot resistance in Experiment 9A (Table 6). Progeny families from two USDA-ARS Fort Collins populations and one USDA-ARS East Lansing mapping population were evaluated in experiment 10A (Table 7). Breeding lines and family progeny were also tested at the BSDF Nursery in Kimberly, ID (Table 8).

Cercospora Leaf Spot/Curly Top Resistant (LSR/CTR) Breeding Populations Currently under Development.

- 1. Cercospora leaf spot and curly top resistant monogerm base population from a polycross of FC607 and FC604 with two Salinas germplasms 2859 and 2890.
 - C. 2890 (sp) = 0790 mm aa x 1890 (Salinas); is seed from aa plants open pollinated by Aplants. 0790 = population-790 cycle 5 synthetic by S₁ progeny, aa, mm, O-type, good combining ability, adapted to California, S^f. 1890 = BC population to population 790 to get Rz equivalent, remains variable for M-:mm, Rz-:rzrz, etc.
 - D. 2859 m (sp) = 1859, 1859R aa x A- (Salinas); Released in 1992 as C859. Sf, similar to 2890, but should have higher curly top resistance. Segregates and variable for M-:mm, Rz-:rzrz, A-:aa, predominant background is lines like C563.
- 2. Cercospora leaf spot and curly top resistant multigerm base population from a polycross of FC902 with two Salinas germplasms 278 and 4918.
 - A. 278 (Iso 83) = RZM R078; R278 is Rz (segregates Rz--:rzrz) version of C46. It should be S^sS^s, MM.

- B. 4918 (sp) = RZM 3918aa X A-, 142 aa plants; This is an increase of released material C918. It should be Multigerm, over 75% Sf and segregating for A-, R-, Rz-, VY, CT, Erw, & PM.
- 3. Cercospora leaf spot and curly top resistant multigerm, self-incompatible base population from a polycross of FC607 x [SR87, MonoHy A4, MonoHy T6, & MonoHy T7]
- 4. Seed from FC709-2 x FC907 was sent to Larry Campbell at Fargo to cross to Sugar beet root maggot resistant germplasm to develop a population that will produce pollinators with resistance to Rhizoctonia, Cercospora, and Root maggot.
- 5. Two tetraploid pollinators (FC6064X and FC6074X) were crossed to a high sucrose tetraploid population in order to produce a tetraploid Cercospora resistant pollinator population with better combining ability.

Progress in 1999

Advanced breeding lines of *Cercospora* resistant germplasms were evaluated in the ARS leaf spot nursery at Ft. Collins. These lines are part of the resistant germplasm development effort in which a new germplasm should be released from the "pipeline" every two to four years. The above populations currently are in different stages of development.

- 1. Selections were made 1998 among half-sib progeny rows of the monogerm population. Families were selected based on leaf spot resistance, curly top resistance, and combined leaf spot and curly top resistance. They were increased and will be tested in the Cercospora nursery and curly top nursery in 2000. They have been also planted in Salinas to select for the single gene source of Rhizomania resistance. Selected roots are being recombined and the resulting population(s), tested, O-type screened, released, or reselected.
- 2. Plants (F₂) from the CTR/LSR multigerm cross (2) were planted in the breeding nursery last summer and *aa* females crossed to the (FC709-2 x FC907)F₂ roots selected in the Rhizoctonia nursery. This seed has been bulk increase and the resulting population will be a source of curly top resistant multigerm pollinators with leaf spot and Rhizomania resistance.
- 3. Plants (F₂) from the Fort Collins and Fargo joint project (3) were grown in the breeding nursery and these roots were planted in Masonville selfed, taking advantage of the 'pseudo self-fertility' that occurs in this environment. This selfed seed was progeny tested in 1999. The most resistant families will be recombined and selected for yield factors. This population will be a source of highly leaf spot resistant multigerm pollinators with curly top resistance and good combining ability for agronomic traits.
- 4. Plants (F_1) from this multigerm cross (4) have been grown in the greenhouse and selfed to produce F_2 seed.
- 5. Bulked F₂ seed was planted in the Rhizoctonia and curly top nursery and half-sib families in the Cercospora nursery. The F₁ has been bulk increased and F₂ seed will be planted in the 2000

Cercospora nursery to select for sucrose and resistance to Cercospora leaf spot.

The seed from the above mentioned populations will be developed and advanced after testing. Development of a resistant germplasm line generally takes 7 years. A longer time may be necessary to incorporate multiple disease resistances. In an established program, a "pipeline" of lines in various stages of development and evaluation is the norm. Hence, the release of new germplasm usually occurs every 2 to 4 years.

Genetic information developed in this research will be used to execute additional cycles of pathogen inoculation, plant selection, and recombination among germplasms that we have in our leaf spot improvement program. Results of these tests will be the basis of decisions about specific germplasm, i.e., retain, discard, recombine, release, etc. Germplasms likely to be useful for variety improvement are identified and released for use by other sugar beet breeders. Breeding techniques are compared in developing these germplasm and information on the efficacy and efficiency of these techniques generated.

Table 5. Experiment 7A, 1999. Leaf Spot Evaluation of USDA-ARS Fort Collins, Salinas, East Lansing, and Fargo breeding lines.

				Disease Index ¹	
Entry	Identification		September 7th	September 14th	September 22nd
		LSD _{0.05}	0.77	0.93	1.02
LSS 2 (931002)			5.0	5.2	6.5
٠	2)		2.7	2.8	3.3
تق			3.6	3.9	4.7
99102-00	99302-00	East Lansing - JS	2.7	2.8	3.0
98J02x05	98J02x05	East Lansing - JS	2.8	2.8	3.3
99125-023	99J25-023	East Lansing - JS	2.7	2.8	3.5
99119-00	99119-00	East Lansing - JS	2.8	3.0	3.7
EL 51	WC9800435L	East Lansing - JS	2.7	3.0	3.8
99131-00	99131-00	East Lansing - JS	3.0	3.3	3.8
WC980437	WC980437	East Lansing - JS	3.0	3.7	4.0
99133-00	99133-00	East Lansing - JS	4.0	4.0	4.7
98J28-02	98J28-02	East Lansing - JS	3.8	4.2	4.8
EL 38	WC980433	East Lansing - JS	4.3	4.5	5.7
96N0012	Low Sodium	Fargo - LC	2.8	3.3	3.7
96N0011	Low Potassium	Fargo - LC	3.3	3.3	4.3
97N0132	F1015	Fargo - LC	4.2	4.7	5.5
6000N96	Low amino-N	Fargo - LC	3.8	4.2	5.8
98N0057	F1016	Fargo - LC	3.8	4.3	6.0
B-5931	Commercial	Fargo - LC	3.2	3.2	3.5
75 (3712)/25 (5931)) Commercial	Fargo - LC	3.8	4.2	4.8
25 (3712)/75 (5931) Commercial) Commercial	Fargo - LC	3.7	3.8	5.0
50 (3712)/50 (5931)) Commercial	Fargo - LC	3.8	4.0	5.2
B-3712	Commercial	Fargo - LC	5.3	5.8	6.0
97-SP22-0	Inc. SP7622-0 (LSR ck) - Iso 86	Salinas - RL	3.3	3.7	4.2
Monodono	(resistant check) - HM	Salinas - RL	3.5	3.7	4.3
EL-02	Rzm EL (Rz x sm. root) - Iso 53	Salinas - RL	3.8	4.2	4.7
Ippolita	(resistant check) - HM	Salinas - RL	3.5	3.7	4.7
CR811	Rzm 711, CR09/10 - Iso 86	Salinas - RL	3.5	3.7	4.7

Table 5. Experiment 7A, 1999. Leaf Spot Evaluation of USDA-ARS Fort Collins, Salinas, East Lansing, and Fargo breeding lines.

				Disease Index ¹	
Entry	Identification		September 7th	September 14th	September 22nd
		LSD _{0.05}	0.77	0.93	1.02
LSS ² (931002)			5.0	5.2	6.5
	1(H2)		2.7	2.8	3.3
Trial Mean			3.6	3.9	4.7
US H11	LSS check - HH	Salinas - RL	3.8	3,8	8.4
K869	Rzm Y769, C69 - Iso 9	Salinas - RL	3.7	4.3	5.0
EL-04	Rzm EL (Rz x sm. root) - Iso 54	Salinas - RL	3.7	4.2	5.0
CR812	Rzm 712 - Iso 87	Salinas - RL	3.8	4.0	5.3
R827	Rzm R727A, B - Iso 12	Salinas - RL	4.5	5.3	5.3
CR813	Rzm 713 - Iso 88	Salinas - RL	3.5	4.2	5.5
Y875	Rzm 775 - Iso 11	Salinas - RL	4.3	5.0	5.5
R726	Rzm-ER R526, C26 - Iso 66	Salinas - RL	3.5	4.5	5.8
8932M(CTR)	7932 CT,aaxA - Sp 12	Salinas - RL	3.8	4.5	6.3
Rifle	Commercial Check - SS	Salinas - RL	5.5	6.2	6.5
B4430R	L4430 (LSS ck)	Salinas - RL	7.3	7.0	7.7
911026HO	FC715	Fort Collins	3.0	2.8	3.3
831085HO	FC708	Fort Collins	3.0	3.0	3.5
97A050	FC607	Fort Collins	3.0	3.0	3.7
921021	FC703-5	Fort Collins	3.0	3.5	3.8
921024	FC709-2	Fort Collins	3.2	3.5	4.0
921025	FC728	Fort Collins	3.5	3.5	4.0
921022	FC702-7	Fort Collins	2.8	3.2	4.3
951017	FC727	Fort Collins	3.3	4.0	4.7
911031	FC717	Fort Collins	3.5	4.2	4.8
¹ Disease Index	¹ Disease Index is based on a scale of 0 (=healthy) to 10 (=dead)	=dead).			
$ ^2$ The Leafspot S	² The Leafspot Susceptible Check is SP351069-0.				
The Leafspot F	³ The Leafspot Resistant Check is ((FC504CMS x FC502/	CMS x FC502/2) x SP6322-0).			

Table 6. Experiment 9A, 1999. Leaf Spot Evaluation of USDA-ARS Fort Collins breeding lines.

				Disease Index ¹	
Entry	pI	Identification	September 7th	September 14th	September 22nd
	-	$LSD_{0.05}$	0.77	0.92	1.02
LSS 2 (931002)		进 一	5.0	5.7	6.3
	2)		2.3	2.0	2.7
<u>Te</u>	2,-		3.2	3.3	3.9
911026HO	FC715	released	2.3	2.3	2.7
99A003	EL 52	released	2.3	2.5	2.8
921024	FC709-2	released	2.5	2.7	2.8
961013HO	FC506	released	2.8	3.0	3.0
96A009	EL 50	released	2.7	2.3	3.0
831085HO	FC708	released	3.0	2.5	3.0
99A001	892016H2	FC607 OT/Beta 2007 (2X)	3.0	2.7	3.3
921022	FC702-7	+ 7 cycles Rhizoc	2.8	2.8	3.3
97A050	FC607 released	ed	2.8	2.5	3.3
98A152	892010H2	FC607 OT/ Hilleshög 8277	3.0	2.7	3.3
971017	FC710 (4X)		2.8	3.0	3.3
971018	FC712 (4X)		2.5	2.8	3.5
921021	FC703-5	released	3.0	3.0	3.7
861039	FC712	released	3.5	3.2	3.7
961010HO1	FC722CMS	C718/FC708 CMS	3.2	3.3	3.7
961010HO	FC722-1	C718/FC708	3.3	3.0	3.7
951017	FC727	released	3.5	3,3	3.7
961015	FC720-1	C718//(C718/FC708)	3.0	3.0	3.7
961011HO1	FC607/FC708CMS	8CMS	3.3	3.5	3.8
991014	Rhizoctonia F	Rhizoctonia Resistant Multigerm pop (2915/FC709-2)	3.2	3.2	3.8
951014	(2890aa & 28	(2890aa & 2859aa) x FC708	3.3	3.5	3.8
99A006	SR 94	released	3.3	3.5	4.0
971020	FC907-1	FC607/FC701 BC4	3.5	3.5	4.2

Table 6. Experiment 9A, 1999. Leaf Spot Evaluation of USDA-ARS Fort Collins breeding lines.

					Disease Index	1		
Entry		Identification		September 7th	 September 14th		September 22nd	p
			LSD _{0.05}	0.77	0.92		1.02	
LSS ² (931002)				5.0	5.7	3	6.3	
LSR ³ (821051H2)	2)			2,3	2.0		2.7	
Trial Mean			1 0 1	3.2	3.3	-	3.9	
911031	FC717	released		3.3	3.5		4.2	
921025	FC728	released		3.3	3.5		4.3	
99A002	892012H2			3.7	4.5		4.7	
961012HO	FC712/MonoHy A4	oHy A4		3.7	4.3		8.4	
961011HO	FC607/FC708	80		3.8	4.2		5.0	-
961012HO1	FC712/MonoHy A4 -	oHy A4 - CMS equivalent		4.0	4.3		5.0	
951016HO1	FC723CMS	FC723CMS EL44/FC708 CMS		3.5	4.0		5.2	
Red Beet Filler				4.2	5.2		5.7	
951016HO	FC723 EL44/FC708 1	4/FC708 mm		4.3	4.5		5.8	
¹ Disease Index is ba	sed on a scale	¹ Disease Index is based on a scale of 0 (=healthy) to 10 (=dead)	ead).					Г
² The Leafspot Susceptible Check is SP351069-0.	eptible Check i	s SP351069-0.						
³ The Leafspot Resi	stant Check is (³ The Leafspot Resistant Check is ((FC504CMS x FC502/2) x SP6322-0)	x SP6322-0).					

Table 7. Experiment 10A, 1999. Leaf Spot Evaluation of USDA-ARS Fort Collins 1/2 sib and selfed Progeny.

		Disease Index	
Entry	September 7th	September 14th	September 22nd
LSS ² (931002)	5.0	5.7	5.0
LSR 3 (821051H2)	2.3	2.0	2.0
Te	2.7	3.2	3,8
$991004 - 37 961023 \otimes = (FC907 \times FC709 - 2)F2 - Rhzc blk$	2.00	2.50	2.25
$991004 - 25 961023 \otimes = (FC907 \times FC709 - 2)F2 - Rhzc blk$	2.00	2.00	2.50
$991004 - 8 961023 \otimes = (FC907 \times FC709-2)F2-Rhzc blk$	2.25	2.75	2.75
$991004 - 44 961023 = (FC907 \times FC709-2)F2-Rhzc blk$	2.25	2.50	2.75
$991004 - 45\ 961023 \otimes = (FC907 \times FC709-2)F2-Rhzc blk$	2.50	3.00	3.00
$991004 - 35 961023 \otimes = (FC907 \times FC709 - 2)F2 - Rhzc blk$	2.00	2.75	3.00
$991004 - 13 961023 \otimes = (FC907 \times FC709-2)F2-Rhzc blk$	2.25	2.25	3.00
$991004 - 30 961023 \otimes = (FC907 \times FC709 - 2)F2 - Rhzc blk$	2.25	2.25	3.00
$991004 - 17961023 \otimes = (FC907 \times FC709-2)F2-Rhzc blk$	2.00	2.50	3.00
$991004 - 36961023 = (FC907 \times FC709-2)F2-Rhzc blk$	3.00	3.00	3.00
$991004 - 12\ 961023 \otimes = (FC907 \times FC709 - 2)F2 - Rhzc blk$	2.50	3.00	3.25
$991004 - 20 961023 \otimes = (FC907 \times FC709 - 2)F2 - Rhzc blk$	2.25	3.00	3.25
$991004 - 52 961023 \otimes = (FC907 \times FC709-2)F2-Rhzc blk$	2.50	3.00	3.25
$991004 - 7 961023 \otimes = (FC907 \times FC709 - 2)F2 - Rhzc blk$	3.00	3.00	3.50
$991004 - 48 961023 \otimes = (FC907 \times FC709-2)F2-Rhzc blk$	2.25	3.00	3.50
$991004 - 3 961023 \otimes = (FC907 \times FC709 - 2)F2 - Rhzc blk$	2.50	3.00	3.50
$991004 - 38 961023 = (FC907 \times FC709-2)F2$ -Rhzc blk	2.75	3.00	3.50
$991004 - 1 961023 = (FC907 \times FC709-2)F2-Rhzc blk$	2.25	3.00	3.75
$991004 - 15 961023 \otimes = (FC907 \times FC709-2)F2-Rhzc blk$	3.00	3.50	4.00
991004 -18 9610238 = (FC907 x FC709-2)F2-Rhzc blk	3.00	3.50	4.00
$991004 - 4 961023 \otimes = (FC907 \times FC709-2)F2$ -Rhzc blk	3.00	3.50	4.00
991004 -23 9610238 = (FC907 x FC709-2)F2-Rhzc blk	3.25	3.75	4.00
991004 -42 9610238 = (FC907 x FC709-2)F2-Rhzc blk	3.00	3.00	4.00
991004 -22 961023 = (FC907 x FC709-2)F2-Rhzc blk	2.75	3.50	4.25

Table 7. Experiment 10A, 1999. Leaf Spot Evaluation of USDA-ARS Fort Collins 1/2 sib and selfed Progeny.

		Disease Index ¹	
Entry Identification	September 7th	September 14th	September 22nd
LSS ² (931002)	2.0	5.7	5.0
LSR ³ (821051H2)	W. t.	2.0	2.0
991004 -46 9610238 = (FC907 x FC709-2)F2-Rhzc blk	3.00	3.00	4.25
991004 -53 961023 = (FC907 x FC709-2)F2-Rhzc blk	2.75	3,75	4.25
$991004 - 26961023 = (FC907 \times FC709-2)F2-Rhzc blk$	3.00	3.50	4.50
991004 -6 961023 = (FC907 x FC709-2)F2-Rhzc blk	3.00	5.00	4.50
$991004 - 31 961023 \otimes = (FC907 \times FC709 - 2)F2 - Rhzc blk$	3.50	4.50	4.75
$991004 - 27 961023 \otimes = (FC907 \times FC709 - 2)F2 - Rhzc blk$	3.00	3.50	4.75
$991004 - 39961023 \otimes = (FC907 \times FC709 - 2)F2 - Rhzc blk$	3.00	3.50	5.00
991004 -29 9610238 = (FC907 x FC709-2)F2-Rhzc blk	3.50	4.50	5.25
991004H 961023® = (FC907 x FC709-2)F2-Rhzc blk [15 plants]	3.00	3.00	3.25
981028 -11 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.50	2.50	2.50
981028 -36 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.50	2.25	2.75
981028 -23 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.75	2.75	2.75
981028 -21 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	3.00	2.75	3.00
981028 -15 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.25	2.25	3.00
981028 -67 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.25	2.75	3.00
981028 -20 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.50	3.00	3.00
981028 -3 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.50	2.75	3.25
981028 -9 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.75	3.00	3.25
981028 -14 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	3.00	2.50	3.25
981028 -69 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.25	2.25	3.25
981028 -24 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.25	2.25	3.25
981028 -28 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.75	3.00	3.25
981028 -25 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.25	2.75	3.25
981028 -77 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.75	2.75	3.50
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Table 7. Experiment 10A, 1999. Leaf Spot Evaluation of USDA-ARS Fort Collins 1/2 sib and selfed Progeny.

		Disease Index ¹	
Entry	September 7th	September 14th	September 22nd
(931002)	5.0	5.7	5.0
(821051H2)	2.3	2.0	2.0
$\overline{}$	2.7	3.2	3.8
981028 -85 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.25	2.50	3.50
981028 -66 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.50	3.25	3.50
981028 -50 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.75	3.00	3.50
981028 -30 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.75	3.25	3.50
981028 -19 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.75	2.75	3.50
981028 -49 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	3.00	3.25	3.50
981028 -59 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	3.00	3.00	3.50
981028 -61 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.50	3.00	3.50
981028 -71 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.50	3.00	3.50
981028 -57 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.75	3.25	3.50
981028 -53 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.75	3.25	3.75
981028 -7 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.25	2.50	3.75
981028 -51 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.75	3.00	3.75
981028 -2 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.75	3.25	3.75
981028 -1 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.35	3.00	3.75
981028 -55 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.75	3.25	4.00
981028 -75 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	2.75	3.25	4.00
981028 -5 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	3.00	3.75	4.00
981028 -12 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	3.00	3.75	4.00
981028 -13 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	3.50	4.00	4.50
981028 -78 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss7)F2 - blk(Ss7)F3 - Sx	3.00	4.25	4.50
981028 -84 (FC607 x (MonoHy-T6, -A7, -A4, & SR87) - blk(Ss?)F2 - blk(Ss?)F3 - Sx	3.00	4.00	00.9
99EL 01 Mapping Population for M McGrath - USDA-ARS, E Lansing	3,00	3.75	4.25
99EL 02 Mapping Population for M. McGrath - USDA-ARS, E. Lansing	3.50	4.50	5.25

Table 7. Experiment 10A, 1999. Leaf Spot Evaluation of USDA-ARS Fort Collins 1/2 sib and selfed Progeny.

				Disease Index ¹	
Entry	Ţ	Identification	September 7th	September 14th September 22nd	tember 22nd
LSS 2	6	LSS 2 (931002)	5.0	5.7	5.0
LSR 3	ڪ	(821051H2)	2.3	2.0	2.0
Trial Mean	Mea		2.7	3.2	3.8
99EL	04	99EL 04 Mapping Population for M. McGrath - USDA-ARS, E. Lansing	2.75	3.25	4.50
99EL		05 Mapping Population for M. McGrath - USDA-ARS, E. Lansing	3.00	4.00	4.25
99EL	07	Mapping Population for M. McGrath - USDA-ARS, E. Lansing	3.00	3.75	5.00
99EL	08	Mapping Population for M. McGrath - USDA-ARS, E. Lansing	3.00	3.25	4.00
99EL	60	Mapping Population for M. McGrath - USDA-ARS, E. Lansing	2.25	2.25	3.00
99EL	10	Mapping Population for M. McGrath - USDA-ARS, E. Lansing	3.25	4.25	4.50
99EL	11	Mapping Population for M. McGrath - USDA-ARS, E. Lansing	3.50	4.25	5.75
99EL		12 Mapping Population for M. McGrath - USDA-ARS, E. Lansing	3.00	4.00	5.00
99EL 15 6869	15	6989	3.25	4.25	5.50
¹ Disea	se In	¹ Disease Index is based on a scale of 0 (=healthy) to 10 (=dead).			
The I	eafs	² The Leafspot Susceptible Check is SP351069-0.			-
³ The L	eafs	³ The Leafspot Resistant Check is ((FC504CMS x FC502/2) x SP6322-0).			

Table 8. 1999 Curly Top Nursery in Kimberly Idaho.

		Disease	e Index ¹
Seed Source	Description	08/31/98	09/22/98
911032	FC718 - Susceptible Check	5.0	6.0
94A068	Beta G6040 - Resistant Check	2.3	3.7
98A101		3.0	4.7
98A102		3.7	5.3
98A103		3.0	5.0
98A104		3.3	5.0
98A105		4.7	6.3
98A106		4.7	6.0
98A107		4.7	5.7
98A108		4.3	5.3
98A109		3.0	5.7
98A110		3.0	4.7
98A111		3.7	4.7
98A112		3.3	4.7
98A113		4.3	6.0
98A114		4.7	5.7
98A115		3.7	5.0
98A116		3.3	5.3
98A117		3.0	5.0
98A118		3.0	5.3
98A119		4.0	5.3
98A120		5.0	6.3
98A121		5.0	6.0
98A122		5.0	7.0
98A123		4.3	6.0
98A124		5.3	6.0
98A125		4.0	5.0
98A126		4.7	5.7
98A127		3.7	4.7
98A128		3.3	4.7
98A129		4.0	5.7
98A130		5.3	6.3
98A131		6.3	7.7
98A132		4.3	6.3
98A133		3.7	5.7
98A134		3.7	4.7
98A135		3.7	4.7

Table 8. 1999 Curly Top Nursery in Kimberly Idaho.

		Disease	Disease Index ¹		
Seed Source	Description	08/31/98	09/22/98		
911032	FC718 - Susceptible Check	5.0	6.0		
94A068	Beta G6040 - Resistant Check	2.3	3.7		
98A136		3.3	4.7		
98A137		5.0	6.0		
98A138		4.0	5.3		
98A139		4.3	5.7		
98A140		4.3	5.7		
98A141		4.7	5.7		
98A142		5.0	5.7		
98A143		4.0	4.7		
98A057		3.3	5.0		
98A096		3.7	5.0		
98A077		3.7	5.3		
971017	FC710 (4X)	3.7	4.3		
97A050	FC607 released	2.7	4.3		
961011 HO	FC607/FC708	3.3	4.7		
971020	FC907-1 FC607/FC701 BC ₄	3.0	4.7		
961011HO1	FC607/FC708CMS	3.7	4.7		
921024	FC709-2 released	3.7	4.7		
951014	(2890aa & 2859aa) x FC708	3.0	5.0		
991003H	CTR/LSRmm	4.0	5.0		
921021	FC703-5 released	4.0	5.0		
99A002	892012H2	2.7	5.3		
991001	RhzcRmmpop; FC708 & 2890,2859 (Salinas)	4.3	5.3		
921022	FC702-7 + 7 cycles Rhizoc	4.3	5.3		
861039	FC712 released	4.0	5.3		
971018	FC712 (4X) released	4.3	5.3		
991014	Rhizoctonia Res. Multigerm pop (2915/FC709-2)	3.3	5.3		
96101 2HO 1	FC712/MonoHy A4 - CMS equivalent	3.7	5.3		
981037	LSR/CTR/Sucrose	4.0	5.3		
991002PF	RhzcR/LSR/MM/Hspop: 3859, 4918, 278; FC907; FC709-2, FC902; MonoHy-T6,A7,& A4; SR87	4.0	5.3		
99 A 006	SR 94 released	4.0	5.3		
961012HO	FC712/MonoHy A4	4.7	5.7		
95101 6HO 1	FC723CMS EL44/FC708 CMS	3.7	5.7		
961010 HO 1	FC722CMS C718/FC708 CMS	4.3	5.7		
831085HO	FC708 released	5.0	5.7		

Table 8. 1999 Curly Top Nursery in Kimberly Idaho.

			Disease	Index ¹
Seed Source		Description	08/31/98	09/22/98
911032	FC718 -	Susceptible Check	5.0	6.0
94A068	Beta G604	10 – Resistant Check	2.3	3.7
99A001	892016H2	FC607 OT/Beta 2007 (2X)	4.3	5.7
961013HO	FC506	released	4.3	5.7
921025	FC728	released	4.3	6.0
961015	FC720	C718//(C718/FC708)	4.3	6.0
961010 HO	FC722	C718/FC708	5.0	6.0
99A003	EL 52		4.3	6.0
911026HO	FC715	released	4.7	6.0
951017	FC727	released	4.7	6.3
951016HO	FC723	EL44/FC708 mm	5.0	6.3
911031	FC717	released	5.3	7.0
¹ Disease Index i	s based on a s	cale of 1 (=healthy) to 9 (=dead).		

PRE-BREEDING: THE INTROGRESSION OF NEW SOURCES OF CERCOSPORA LEAF SPOT RESISTANCE FROM BETA VULGARIS SPP. MARITIMA AND OTHER EXOTIC SOURCES INTO SUGAR BEET-TYPE POPULATIONS. (BSDF Project 443) L. Panella

A major emphasis of the research mission of the USDA-ARS plant scientists is the collection, documentation, characterization, evaluation, regeneration (maintenance), distribution, and utilization of plant germplasm, especially Plant Introduction (PI) accessions in the USDA-ARS National Plant Germplasm System (NPGS). The Sugar Beet Research Unit at Fort Collins is coordinating the national program for *Beta* germplasm evaluation. In addition to the evaluation for Rhizoctonia and Cercospora resistance, it is crucial that the ARS scientist be involved in the long rang, high risk research problems involved in sugar beet 'germplasm enhancement' or 'pre-breeding'. This is an important component in the overall sugar beet improvement effort of the Fort Collins Sugar Beet Research Unit.

Justification for Research: Cercospora leaf spot (caused by the fungus Cercospora beticola Sacc.) is one of the most widespread diseases of sugar beet and is a serious problem in many sugar beet production areas throughout the U.S. The disease damages the leaves, which, consequently, reduces root yield, percent sucrose of roots, and purity of the extracted juice. Cercospora leaf spot currently is controlled by combining spraying with commercial fungicides and the use of disease tolerant germplasm. The development of Cercospora leaf spot resistant sugar beet lines and hybrids with greater levels of host-plant resistance offers a more sustainable solution to this disease problem.

If the level of resistance available in some Cercospora-resistant experimental breeding lines were present in commercial hybrids (along with good sugar and seed yield), the need for fungicides could be greatly reduced. That continued improvement in genetic resistance to this serious pathogen is still needed is evident by the occurrence of *Cercospora* strains that are tolerant to our most potent fungicides. Additionally, some fungicides may be removed from the market because of their perceived or real threat to the environment.

Finally, the genepool for resistance to Cercospora leaf spot is extremely narrow. Many of the resistant lines are highly inbred, therefore, closely related to one another, and stem from germplasm coming out of Italy in the early 1900s. In the germplasm developed at Fort Collins, continued inbreeding has increased the level of disease resistance, but at the cost of plant vigor. Over the long term, a secure, sustainable response to this disease requires commercial quality hybrids with good host-plant resistance.

Objectives:

- 1. The formation of long range breeding populations through the introgression of Cercospora resistant germplasm from "exotic" sources (*Beta vulgaris* spp. *maritima*, fodder beet, foreign sugar beet landraces from the PI collection, etc.).
- 2. The development of germplasm populations from these long range populations that are of sufficient agronomic quality to be of use to commercial breeders. They will be a source of

leaf spot resistance with differing genetic backgrounds.

3. The development of techniques (both traditional and molecular) to more efficiently introgress the exotic germplasm into sugar beet breeding populations.

Research Progress 1999:

Crosses have been made or are being attempted in the greenhouse on the list of accession below, all of which have been identified as having Cercospora resistance. F_2 is being planted from the F_1 populations, where sufficient seed is available. F_2 seed of three crosses (96A011, 96A015, and 96A016 as donor parents) has been bulk increased in the greenhouse and this is being planted to produce F_3 populations. All three show some biennial plants in our environment and were crossed to genetic male sterile (aa) sugar beets. These F_1 increases should be completed by the beginning of 2001. We are considering re-crossing some of those from which we obtained insufficient F_1 seed, but will concentrate primarily with those populations from which we have sufficient seed.

Plants from those populations producing some biennial plants are being vernalized for 90 days and the populations are being increased (i.e., random mated using the genetic male sterility where possible). The annuals will be handled in a similar fashion once the F_1 populations have been increased. All will be cycled through at least three cycles of random mating.

Accession Number	Donor Designation	Name or Origin	% Bolting without induction 1996 Fort Collins	F ₁ Population	F ₂ Population
96A010	PI 535826	Giant Poly	20%	9 7 1021H2	981031 F ₃ =991026
96A011	PI 535833	Saturn	0%	unsuccessful	
96A014	PI 540593	WB 847	0%	971023H2	
96A015	PI 540596	WB 850	70%	971024H2	981032
96A017	PI 540605	WB 859	25%	971025H2	
96A012	PI 535843	PN MONO 1	100%	971026H21	
96A013	PI 540575	WB 829	100%	971027H2 ²	
96A016	PI 540599	WB 853	50%	971028H2	981033
94A079	#32375 (B. v. ssp. maritima)	Greece	annual	971029H2	
94A080	#36538 (B. v. ssp. maritima)	Greece	annual	971030H2 ³	
94A081	#45511 (B. v. ssp. maritima)	Greece	annual		
94A082	#45516 (B. v. ssp. maritima)	Greece	annual	981002H3	
94A083	#48810 (B. v. ssp. maritima)	Tunisia	annual		
94A084	#48819 (B. v. ssp. maritima)	Tunisia	annual	981004H2	
94A085	#51430 (B. v. ssp. maritima)	Greece	annual	981005H3	

¹Only 16 seed balls produced.

²Only 10 seed balls produced.

³Only 60 seed balls produced.

Development of a resistant germplasm line generally takes seven years. A longer time will be necessary to incorporate disease resistance from more exotic sources. Because this is a new program it will take time for the first germplasm to make it through the process. Once that happens, there will be a "pipeline" of germplasm in various stages of development and the release of new germplasm will occur every two to four years. The incorporation of exotic sources into agronomically acceptable germplasm is a long term proposition - results will not appear overnight. This is the type of long-term, high risk germplasm research that ARS is well-suited to perform.

Materials and Methods: Artificial inoculation with Cercospora beticola and leaf spot scoring will be used to identify the resistant germplasm sources and make selections in the developing populations. The exotic materials will be crossed into sugar beet populations that have been selected for agronomic quality (recoverable sucrose yield). These are currently under development using germplasm received from commercial breeding programs, public sources (e.g., L19), and some high sucrose germplasm from Poland. These sugar beet populations will be self-fertile (Sf) and segregating for nuclear male sterility (A-:aa). Populations will be handled in the following manner: 1) Following the initial cross, a population will be random mated (using aa females because of the self-fertility) for three to four generations to break up linkage groups and remove annual plants. 2) Sugar beet-type mother roots will be selected, selfed, and progeny tested for agronomic performance and disease resistance. 3) Selected roots will be recombined (and backcrossed if desirable) and re-selected until they ready for release. Molecular markers (RFLPs, RAPDs, SSRs, AFLPs, etc.) will be used to expedite the backcrossing program and to follow the change in allele frequencies in the selected populations. Advanced populations will be released to the sugar beet seed industry.

Summary of Literature: Cercospora leaf spot has been an intermittent problem in sugar beet growing areas of the United States where the summers can be hot and humid (Red River Valley, Michigan, Ohio, and, less often, Great Plains growing areas and California). It has been estimated that a severe epidemic can cause up to a 42% loss of gross sugar (Smith and Martin, 1978; Smith and Ruppel, 1973), or up to a 43% relative dollar loss (Shane and Teng, 1992).

Resistance to Cercospora leaf spot has long been a goal of the USDA-ARS sugar beet research program at Fort Collins and researchers there developed the techniques necessary to manage the screening nurseries in such a way as to promote the development of the disease (Ruppel and Gaskill, 1971). A careful crop rotation (sugar beet-barley-barley-barley-sugar beet) and the arid climate and low relative humidity have allowed this to be done in such a manner that there are rarely high enough levels of any other disease present in the leaf spot nursery to confound the results.

There are an estimated 4 or 5 genes responsible for *Cercospora* resistance (Smith and Gaskill, 1970) and broad-sense heritability estimates ranged from 12 to 71% (Bilgen et al., 1969). Narrow-sense heritability estimates of about 24% compared well with realized heritability values, and 44 to 62% of the variation was due environment in this test (Smith and Ruppel, 1974). The large environmental variation has made it difficult to make progress in developing *Cercospora* resistance through mass selection. Incorporation of high levels of leaf spot resistance into varieties with superior agronomic performance also is difficult (Smith and Campbell, 1996) and, therefore, commercial resistant varieties require some fungicide application to provide adequate levels of protection against Cercospora (Miller et al., 1994).

A major problem in the development of *Cercospora*-resistant sugar beet is the loss of vigor due to the continual inbreeding. Coons (1955) noted this and it has been a concern ever since (McFarlane, 1971). The use of hybrid varieties has ameliorated this problem to some extent, but seed production on the highly inbred O-type males and CMS females still is a problem. This is seen in germplasm from both the FC 500 and FC 600 series developed at Fort Collins.

The USDA-ARS National Plant Germplasm System Beta collection has over 2,000 Plant Introduction (PI) accessions. The germplasm used most often in sugar beet breeding is from Beta vulgaris spp. vulgaris, which includes all of the biennial sugar beet types, or from Beta vulgaris spp. maritima, which contains the closely related wild sea beet and has both annual and biennial types. Germplasm with a biennial flowering habit is easier both to introgress and screen. Beta vulgaris spp. maritima has, nonetheless, been used as a source of resistant germplasm. Much of the Cercospora-resistant germplasm in use today came out of Munerati's program in Italy, in which B. vulgaris spp. maritima was the source of resistance genes (Lewellen, 1992). There have been very few new efforts to locate and incorporate other sources of resistance to Cercospora into this narrow germplasm base.

There is an urgent need to continue to create in our *Cercospora*-resistant germplasm a broader genetic base than we have today. As commercial hybrid parents become more inbred, the germplasm base from which these inbred parents are developed must have the diversity necessary to provide for maximum gain through heterosis. Munerati's success, and the research of others, has shown that it can be done if we have the persistence to do it (Bilgen et al., 1969; Doney, 1993; Lewellen, 1995).

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SUGAR BEET RESEARCH

1999 REPORT

Section C

U. S. D. A., A. R. S., Western Regional Plant Introduction Station Pullman, Washington

Dr. Alan Hodgdon, Beta Curator

This research was supported in part by funds provided through the Beet Sugar Development Foundation (Project 290)

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Status Repor	on the Beta germplasm Collection Activities	
by A.	Hodgdon	C3

Status report on the *Beta* germplasm collection activities at the USDA, ARS, Western Regional Plant Introduction Station To the Beet Sugar Development Foundation Curator: Dr. Alan Hodgdon, 2000

Thirty-nine accessions harvested at W-6 in 1999 have been cleaned and weighed. Of these, thirty-six were increased in the greenhouses. Three of these accessions will have to be redone. Only three accessions were successfully increased in the field due to very bad growing conditions. Twenty-five plots froze during the winter, and several surviving plots did not flower well during the heat of the summer. Fifty-four accessions were started at W-6 in 1999. In all, 110 accessions are in the process of increase. The increase priority list has 448 accession as of the end of 1999. Twelve accessions are being increased by seed companies in the U.S. This help is greatly appreciated. Field increases at W-6 have been a problem with poor plant numbers, poor quality seed, and low seed yield.

In the future we will try artificial vernalization, and then spring plant in at our Pullman site. This could solve the problem of plot freeze-out and provide a cooler weather grow-out site. I am not optimistic about this solution since our growing environment is a poor match with that of the wild beets, particularly *Beta maritima*. With the greenhouse increases, we have had good seed yield and quality, but progress has been slowed by deinduction of flowering especially with the wild beets. We are working on changing the post-vernalization conditions to improve flowering.

SEED GERMINATION

One hundred-five *Beta* seed samples were tested for germination in 1999. The seed lab is using a dry germinator which gives better results. No specific seed germination data is available now. However, no sample had less than 20% germination, and most samples were higher than 50%.

SEED STORAGE ACTIVITY

W-6 distributed 643 samples from the *Beta* collection in 1999, and we acquired thirty- one new accessions. There was no new backup activity for the *Beta* collection.

SUGARBEET RESEARCH

1999 Report

SECTION D

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PUBLICATIONS

Abstract of Papers Presented or Published

CAMPBELL, L.G., G.A. SMITH, J.D. EIDE, AND L.J. SMITH. 1999. *Metarhizium anisopliae* as a biocontrol agent for sugarbeet root maggot. J. Sugar Beet Res. 36(3):55.

Only a few insecticides are available for controlling the sugarbeet root maggot (Tetanops myopaeformis). These could become less effective because of the development of resistant root maggot strains or become unavailable because of environmental concerns. An effective biocontrol agent would provide an alternative and, perhaps, more consistent control method. Laboratory results and a 1995 field trial prompted further testing of the entomopathogenic fungus Metarhizium anisopliae (Metschn.). Metarhizium inoculum was prepared by culturing the fungus on heat-killed barley. The inoculated barley was spread evenly over field plots in the fall proceeding the sugarbeet crop, in the spring prior to planting, or both in the fall and spring. Root yields ranged from 49.5 Mg ha⁻¹ when no insecticide was applied to 59.2 when Lorsban (chlorpyrifos) was used to control maggots. The fall, spring, and fall plus spring applications of *Metarhizium* yielded 51.5, 50.9, and 58.9 Mg ha⁻¹, respectively, at Crookston in 1996. The 1997 trials included the same three Metarhizium treatments with an additional application of Metarhizium in the spring of 1996 (prior to planting barley). Root yields for the Metarhizium treatments ranged from 51.4 to 57.6 Mg ha⁻¹, compared to 57.5 Mg ha⁻¹ when Lorsban was applied and 48.7 Mg ha⁻¹ in the absence of maggot control in 1997. Yield differences between treatments were not significant in 1998 because of reduced root maggot pressure, but appeared to follow the pattern observed in the 1996 and 1997 trials. Results, to date, have been encouraging; however, additional information on application rates and timing, formulations, and the effectiveness of *Metarhizium* in more environments will be required before commercialization is feasible.

CAMPBELL, L.G., A.W. ANDERSON, L.J. SMITH, AND R. DREGSETH. 1999. Root yield losses associated with sugarbeet root maggot damage. J. Sugar Beet Res. 36(1-2):56.

Sugarbeet root maggot, *Tetanops myopaeformis*, is the major insect pest of sugarbeet in Minnesota and Eastern North Dakota. Root maggot damage is routinely rated on a 0 (no damage) to 5 (severe) scale. Forty-two trials were utilized to examine the relationship between visual damage and root yield. The mean damage rating in the absence of insecticides was 3.3, compared to a mean of 1.7 for the highest yielding treatment in each trial. Mean root yield of the highest yielding treatment in each trial was 48.8 Mg ha⁻¹, compared to a mean of 29.0 Mg ha⁻¹ when no insecticides were applied. Regression analyses within individual trials indicated the yield loss associated with each increment of the damage rating scale fluctuated widely, ranging from near zero to 15.7 Mg ha⁻¹. The percent yield reduction in the absence of

insecticides ranged from 9.8% to 83.6% when compared to the treatment providing the most effective control in each test. The regression equation from a combined analysis indicated that little or no yield loss occurs with damage ratings below 1.4. These results are useful in estimating losses, developing recommendations, and providing a standard of comparison for alternative control strategies.

KLOTZ, K.L. 1999. Sucrose metabolizing enzymes and sucrose losses in sugarbeet. Sugarbeet Research and Extension Reports, p.145-147.

Developmental changes in the activities of the major sucrose catabolizing enzymes of sugarbeet roots were determined. In seedling roots, the acid invertases were the predominant sucrolytic enzymes. Soluble and insoluble acid invertase activities were greatest in two week old sugarbeet roots. After two weeks, their activities dropped precipitously to nearly negligible levels. Soluble acid invertase activity was due to a single isoenzyme. Alkaline invertase activity was also greatest in two week old roots. Alkaline invertase, however, was present only at low levels throughout development. Two alkaline invertase isoenzymes were present at all developmental stages, but their relative contribution to total alkaline invertase activity changed with root development. Sucrose synthase was the major sucrose utilizing enzyme in sugarbeet roots six weeks of age or older. Two sucrose synthase isoenzymes contributed to sucrose synthase activity. Only one sucrose synthase isoenzyme was evident during the first twelve weeks of growth. Two sucrose synthase isoenzymes were present after sixteen weeks.

WEILAND, J. J. AND SUNDSBAK, J. L. 2000. Differentiation and detection of sugarbeet fungal pathogens using PCR amplification of actin coding sequences and the ITS region of the rRNA gene. Plant Disease. 84:475-482.

The DNA sequences of the actin genes of several fungi were compared and highly conserved regions in the coding sequence were identified. Deoxyoligonucleotide primers were synthesized based on conserved sequence blocks in the 5' and 3' ends of the open reading frame encoding the actin protein. In addition, primers (ITS1 and ITS4) based on conserved regions of the ribosomal RNA (rRNA) genes of fungi were synthesized. Use of the primers in the polymerase chain reaction (PCR) resulted in the amplification of DNA products from the genomes of sugarbeet fungal pathogens of a size consistent with the amplification of the actin gene and rRNA gene sequences, respectively, in these fungi. With one primer pair (5FWDACT and MIDREVACT) directed to the actin gene, the major products amplified from the DNA of Aphanomyces cochlioides, Pythium ultimum, Cercospora beticola, Phoma betae, Fusarium oxysporum, and Rhizoctonia solani were of the sizes of 0.9, 0.9, 1.1, 1.1, 1.2 and 1.7 kilobasepairs (kbp), respectively, whereas no product was generated from the DNA of sugarbeet (Beta vulgaris L.). Restriction endonuclease digestion of products amplified using 5FWDACT and MIDREVACT permitted the differentiation of A. cochlioides from A. euteiches. Use of ITS1 and ITS4 in PCR reactions employing the same template DNAs and reaction conditions yielded single products of 0.7, 0.8, 0.5, 0.5, 0.6, and 0.7 kbp, respectively, as well as a 0.7 kbp product from DNA of sugarbeet. The data indicate that actin and rRNA gene sequences are appropriate targets for the development of PCR-based strategies for distinguishing sugarbeet fungal pathogens at the genus level. The presence of A. cochlioides DNA in extracts of diseased sugarbeet seedlings was detected using PCR with primers 5FWDACT and MIDREVACT.

WEILAND, J.J. 2000. A survey for the prevalence and distribution of *Cercospora beticola* tolerant to triphenyltin hydroxide and mancozeb and resistant to thiophanate methyl in 1999. 1999 Sugarbeet Research and Education Reports, Cooperative Extension Service, North Dakota State University. 30:236-239.

Triphenyltin hydroxide (TPTH) has been used extensively in the Northern Great Plains in recent years for the control of *Cercospora* leaf spot on sugarbeet. Although mancozeb and, to a lesser extent, the benzimidazole fungicides often are implemented in conjunction with TPTH for optimum leaf spot control, TPTH continues to be the most widely used compound for control of the disease. Testing in our USDA-ARS Fargo laboratory of *Cercospora* that was isolated from leaf spot in the sugarbeet fields in North Dakota and Minnesota for the tolerance or resistance to fungicides first revealed tolerance to TPTH in 1994. The testing program has continued to the present and now includes surveying for tolerance to mancozeb. Testing for baseline tolerance to tetraconazole is also beginning this year, as this represents new chemistry available to the grower for the control of leaf spot disease. As in previous years, fields in the southern Minnesota growing region and in all factory districts from Wahpeton to Drayton in the Red River Valley were surveyed. Samples were tested for resistance to thiophanate methyl (TM; a benzimidazole fungicide) and for tolerance to TPTH and mancozeb at two different exposure levels.

WEILAND, JOHN J., AND ROBERT T. LEWELLEN, J. MITCH MCGRATH, LEE PANELLA, AND MING H. YU. 2000 tagging of disease resistance genes in sugarbeet (beta vulgaris L.) With molecular genetic markers. Abstracts of the Plant and Animal Genome VIII Meeting. p45 of Abstract Book.

Resistance to numerous diseases pests in sugarbeet appear to be conferred by monogenes. These include resistance to powdery mildew, Erwinia vascular necrosis, beet mosaic virus, and Fusarium stalk rot. The inheritance of resistance to the cyst nematode, *Heterodera schachtii*, is monogenic and the inheritance of resistance to the root knot nematode is being evaluated. These pathosystems are being used as models for the generation of molecular genetic markers tagging genes for disease resistance in sugarbeet. Markers generated from the study will be used to evaluate the linkage and location in the sugarbeet genome of genes conferring resistance to several pathogens. In addition, the markers will be useful in the introgression of disease resistance genes into sugarbeet parent lines using marker-assisted selection

and in future cloning and analysis of these genes. The use of resistance gene analog (RGA) sequences is being incorporated into the resistance gene detection strategies. Such sequences may permit the identification of quantitative trait loci that contribute to genetically-complex resistance in sugarbeet to rhizoctonia root rot, Cercospora leaf spot, and aphanomyces black root diseases. The status of a project aimed at tagging a monogene conferring resistance to powdery mildew in sugarbeet caused by *Erysiphe polygoni* DC will be presented.

WEILAND, J. J. AND LEWELLEN, R. T. 1999. Generation of molecular genetic markers associated with resistance to powdery mildew (*Erysiphe polygoni DC*) in sugarbeet (*Beta vulgaris* L.). 9th International Congress on Molecular Plant-Microbe Interactions. July 25-30th, 1999, Amsterdam, The Netherlands.

Powdery mildew caused by Erysiphe polygoni DC can be devastating to sugarbeet production particularly in warm, dry climates. Although resistance to certain races of E. polygoni exists in sugarbeet, powdery mildew disease is typically controlled though the use fungicides. The identification of broad resistance to sugarbeet powdery mildew in the wild beet B. vulgaris spp. maritima was followed by the incorporation of this resistance into sugarbeet by recurrent backcrossing and progeny testing. Germplasm accession C37 was used as the susceptible, recurrent parent and P604 is the F₂BC₃ population at the intermediate stage of the introgression. Three DNA pools each were produced for C37 and P604; each pool was comprised of the DNA from 7 individual plants. A diprimer adaptation of the RAPD analysis was applied to the DNA pools, where one of the primers was composed of a sequence homologous to that encoding a core sequence found in many plant disease resistance genes. Amplified products were identified that were associated with all three DNA pools derived from P604 plants, but with none of the three DNA pools derived from C37. The possibility that some of the amplified products contain sequences of the gene conferring resistance to sugarbeet powdery mildew is discussed.

CHARACTERIZATION OF GENE AND GENE PRODUCTS INVOLVED IN CERCOSPORA RESISTANCE IN SUGARBEET.

Project 601

John J. Weiland

A glucanase enzyme induced in sugarbeet that is infected with *Cercospora beticola* was identified during the course of the project. Using protein sequence data of the enzyme, PCR primers designed for the gene are being used to clone the gene sequences. Once cloned, the sequences can be used as a probe to examine the association of resistance to Cercospora leaf spot disease in sugarbeet populations segregating for this trait.

Elaboration of the approaches outlined in project 601 and application of these approaches to numerous pathogens of sugarbeet are the topic of a new proposal being submitted to the BSDF by J. Weiland ("Mechanisms of resistance in sugarbeet to fungal and bacterial pathogens"). Fundamental to the new project is the use of molecular biology techniques to determine important biochemical players in the defense of sugarbeet from pathogen attack.

Direct biochemical evaluation of the defense process remains an integral component of the analysis. Novel changes in the pattern of isozymes of esterase, acid phosphatase, and peroxidase have been shown to be associated with infection by *C. beticola* (Fig. 1). Those activities that exhibit the most drastic changes as a result of fungal infection will be examined in greater detail using time-course studies. A comparison of the regulation of these activities in leaf spot susceptible and leaf spot resistant germplasm will point to candidate genes underlying the resistance.

By incorporating molecular biology into the analysis of the biochemistry of resistance, novel genes that may confer resistance to sugarbeet through genetic engineering or other means will be more readily obtained. As an example, the cloning of the *sor* (singlet-oxygen resistance) gene from *C. beticola* in our laboratory may find use in engineered sugarbeet for enhancing leaf spot resistance. In addition, a polygalacturonase inhibitor protein (PGIP) gene from *Beta webbiana* has been amplified by PCR and cloned in our lab (Fig. 2). The future tailoring of potential antifungal proteins such as the PGIP gene using recombinant DNA techniques may lead to the development of antifungals with broad spectrum actives against many sugarbeet pathogens.

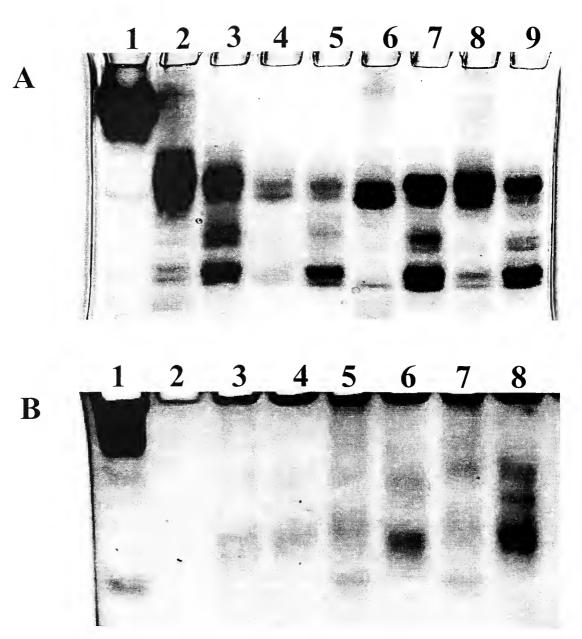


Figure 1. Sugarbeet isozymes of peroxidase (A) and esterase (B) separated by native polyacrylamide gel electrophoresis. In A, even numbered lanes represent extracts of healthy sugarbeet tissue from germplasm of Ultramono (lane 2), BS-S (4), BS-R, (6), and FC607 (8). Extracts from Cercospora lesions on leaves were made and run in lanes 3,5,7,9, representing (in order) the same germplasm sources. Horseradish peroxidase was run in lane 1. The gel was stained with 3-amino-9-ethylcarbazole. For the esterase gel in B, an extract from cultured C. beticola was run in lane 1. Extracts of healthy leaf tissue from sugarbeet Ultramono (lane 8) and FC607 (6) are compared to extracts from Cercospora lesions in lanes 7 and 5, respectively. Lane 4 represents an extract of healthy Ultramono tissue which is compared to the extract in lane 3 from necrotic Ultramono tissue resulting from infiltration with 100 μ M purified cercosporin toxin. Esterase activities were visulized by UV light after treatment of the gel with 4-methylumbelliferyl butyrate. In both A and B, note changes in isozyme pattern after infection with C. beticola.

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pir | S47965 polygalacturonase inhibitor protein - tomato >gi | 469457 (L26529)
                  polygalacturonase inhibitor protein [Lycopersicon
                                    esculentum]
                                    Length = 327
                  Score = 184 bits (462), Expect = 1e-45
 Identities = 113/301 (37%), Positives = 156/301 (51%), Gaps = 31/301 (10%)
Query: 32 CNPQDKKALLEIKHHFHNASAFSNWDPNTDCCSDWFGILGCDSHGR-----ILQLDISS 85
           CNP+DKK LL+IK N ++WDPNTDCC W+ ++ CD
Sbjct: 23 CNPKDKKVLLQIKKDLGNPYHLASWDPNTDCCY-WY-VIKCDRKTNRINALTVFQANISG 80
Query: 86 R------NLTG-IPSSLGQLHKVNTILLNSNNLSGRIPSFFSFMKS 124
                               NLTG IP ++ +L + + L+ NL+G IP F S +K+
Sbjct: 81
          QIPAAVGDLPYLETLEFHHVTNLTGTIPPAIAKLTNLKMLRLSFTNLTGPIPEFLSQLKN 140
Query: 125 LQSY-LYDNQLTGMIPSSLARLPKLLDINLGYNQLTGSIPQXXXXXXXXXXXXXXXXXYYFNK 183
               L NO TG IPSSL++LP LL + L N+LTG+IP+
Sbjct: 141 LTLLELNYNQFTGTIPSSLSQLPNLLAMYLDRNKLTGTIPESFGRFKGPNIPDLYLSHNS 200
Query: 184 LSGPIPRSFGKXXXXXXXXXXXMFTGDASNLFSRDNMELFSIDISSNRFHFDFSKVVLSR 243
           L+G +P S G GD S LF ++ ID+S N FD SK
Sbjct: 201 LTGHVPASLGDLNFSTLDFSRNKLEGDVSFLFGKNKTSQV-IDLSRNLLEFDISKSEFAE 259
Query: 244 KLVYLNVSHNAIYGSLPKNLGQLSLQRIDVSFNQLCGKIPTGRRLKQFSPALFSHNKCLC 303
          L+ L+++HN I+GSLP L + LQ +VS+N+LCG+IP G L+ F
Sbjct: 260 SLISLDLNHNRIFGSLPPGLKDVPLQFFNVSYNRLCGQIPQGGTLQSFDIYSYLHNKCLC 319
Query: 304 G 304
Sbjct: 320 G 320
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Figure 2. A putative polygalacturonase inhibitor protein (PGIP) sequence predicted from the DNA sequence of clone amplified from *Beta webbiana*. Oligonucleotide primers were made based on conserved regions of PGIP genes from other plant species. The amplified DNA from *B. webbiana* was cloned and the sequence obtained by standard methods. BLAST-based search of the Genbank sequence database with the translated DNA sequence revealed similarity to PGIP genes from other plants. The "Query" sequence is that from *B. webbiana*, whereas the "subject" sequence found by the search is from tomato (*Lycopersicon esculentum*).

DEVELOPMENT OF A GREENHOUSE ASSAY FOR RESISTANCE TO RHIZOCTONIA ROOT ROT

Project 610

John J. Weiland and Lee Panella

Methods for the evaluation of sugarbeet for resistance to root rot caused by Rhizoctonia solani AG2-2 presently involve the generation of disease in replicated field plots. The development of a resistance screening method that could be performed in the growth chamber or greenhouse would enable researchers to evaluate candidate breeding lines for levels of resistance before use in test hybrids. In recent years, the ARS lab in Fargo has refined a technique for the inoculation and rating of young roots with *R. solani* AG2-2. A protocol was presented last year that permits roots of test germplasm to be evaluated at 8 weeks post-seeding. Ranking of test germplasm according to levels of disease was similar to that observed for the performance of the accessions in the root rot disease nursery at Fort Collins, CO.

The techniques for inoculation and plant rating are as follows. Briefly, one or two sugarbeet plants are grown in 6" pots to the 5 week-old stage in a greenhouse that is maintained at an average temperature of 25°C and alternating between a 16 hr day period and an 8 hour dark period. Since 50 roots are inoculated per trial, the rearing of at least 60 plants is recommended. Two weeks prior to plant inoculation, *R. solani* AG2-2 is plated onto potato dextrose agar and incubated at 22°C in the dark. One week prior to inoculation, sterile barley grain is sprinkled onto the plated *R. solani* culture and the plates are sealed with Parafilm and returned to the incubator. The barley grains become infested with the fungus within one week. For the inoculation, two infested barley grains are place next to the root surface of a 5 week-old sugarbeet plant at ~2 cm below the surface of the soil. The soil is replaced over the grain inoculum and the plants are watered immediately after all of the plants have been inoculated.

One week after inoculation, plants of a highly susceptible check accession or variety are examined at 3-day intervals in order to monitor disease progress. When greater than 50% of the roots of this accession exhibit severe root rot (>90% of root surface exhibiting rot), all of the roots in the experiment are dug up and rated for root rot severity. This typically occurs at about 14 days post-inoculation. A 0 to 4 scale is used for evaluating root rot severity, where a plant exhibiting no disease is considered a 0 reaction, a root lesion effecting 10% or less of the root surface is a 1 reaction, a root lesion covering 11 – 50% of the root surface is a 2 reaction, root rot covering 51-89% of the root surface is a 3 reaction, and rot on ~90% of the root surface or the plant is dead represents a 4 reaction. By multiplying the data by 7/4, a comparison can be made between the data obtained using the 0-4 scale with that using the 0-7 scale employed at the Fort Collins disease nursery.

In 1999, the technique for evaluating sugarbeet roots for resistance to *R. solani* was applied to a mapping population developed by J.M. McGrath (ARS-East Lansing) and segregating for resistance to Rhizoctonia root rot was evaluated using the greenhouse method. Highly resistant and highly susceptible progeny from the cross will be used to identify molecular genetic markers that co-segregate with root rot resistance. Use of such markers could significantly reduce costs in a breeding program, by substituting marker detection for disease screening. In addition, crosses were initiated between FC403cms, possessing low root rot resistance, and the highly root rot resistant FC709-2,

both produced at the ARS Fort Collins nursery. Interpollination between F1 progeny from this cross will yield F2 progeny varying in resistance to *R. solani*. Highly susceptible and highly resistant F2 progeny will be used for the preparation of DNA and tagging of loci contributing to root rot resistance in sugarbeet using accepted marker methods (amplified fragment length polymorphism, random amplified polymorphic DNA, etc.). Application of DNA marker technology to genetic resistance in sugarbeet to *R. solani* and other fungal pathogens of sugarbeet will be continued in 2000 under a new project sponsored by the BSDF.

POLYMERASE CHAIN REACTION (PCR)-BASED DETECTION OF APHANOMYCES COCHLIOIDES USING ACTIN GENE SEQUENCES.

Project 620

John J. Weiland

A number of soil fungi have the capability to cause disease in sugarbeet and these include *Rhizoctonia solani*, *Aphanomyces cochlioides*, *Pythium aphanidermatum*, *P. ultimum*, and *Fusarium oxysporium*. When seedling damping off or adult plant root rot occur, diagnosis of the causal agent of the disease can be a time-consuming process (days to weeks). Culture of the organisms from an infected area of the sugarbeet root can lead to the recovery of a plethora of fungi, many of which have colonized the infected tissue as saprophytes.

The polymerase chain reaction (PCR) is a DNA based technique for amplifying specific sequences from the genomes of organisms. PCR technology has impacted many fields of biology, including the area of disease diagnosis in both plants and animals. Diagnostics using the PCR are sensitive and highly discriminatory, since they target genome regions whose DNA sequences have diverged throughout evolution. PCR-based diagnostics also require little time for a result to be secured (within one to two days), making them attractive to high-throughput diagnostic laboratories.

The interests in our laboratory include the development of novel diagnostic tools for disease-causing fungi in sugarbeet, as well as the development of tools for investigating the biochemistry of sugarbeet pathogenesis by fungi. For this reason, we designed our PCR assay for the discrimination of sugarbeet fungal pathogens upon DNA sequences of the actin gene. Actin is a protein found in all eukaryotes and the gene coding for actin possesses sequence blocks of both high similarity, as well as of high divergence, across all eukaryotes. This facilitates the design of DNA "primers" that recognize the highly similar sequences in order to detect potential size variation in the actin gene that can be used to "fingerprint" and discriminate one sugarbeet pathogen from another. Actin is also a highly expressed gene and the cloning and re-engineering of actin gene sequences might provide a useful tool for gene transfer studies with sugarbeet fungal pathogens.

In 1999, the results of application of the PCR technique to all major sugarbeet fungal pathogens was summarized (see Weiland and Sundsbak in Publications section). Future work will focus on the design of primers that will permit the robust detection of *A. cochlioides* without amplification of DNA from potentially contaminating DNA from *A. euteiches* or other resident fungi. In addition, observations of DNA polymorphism within the actin gene will be used in conjunction with amplified fragment length polymorphism (AFLP) data in *A. cochlioides* and virulence data in order to assess genetic diversity of *C. beticola* in sugarbeet in the U.S.

THE DEVELOPMENT OF DYNAMIC GENE POOLS FROM BETA MARITIMA SOURCES

Project 630

Larry G. Campbell

Since heterosis generally is enhanced by increasing the genetic diversity of the parents, the introduction of desirable germplasm from previously unused sources is essential to the success of long-range hybrid development programs. Because of its background and the need for specific characteristics such as cytoplasmic male sterility, monogerm, and different disease resistance, the sugarbeet breeding pools are believed to be genetically limited (Lewellen, 1992). Although there appears to be sufficient variability for short term gains, long term progress may very well depend upon the infusion of additional variation into the crop.

Potential sources of genetic variation not now being utilized fully include 1) old land races of sugarbeet, table beet, and fodder beet; 2) new naturally occurring or induced mutations; and 3) wild relatives. New sources of genetic variation should produce fertile offspring when crossed with sugarbeet and be genetically unique and diverse, compared to commercial sugarbeet. Of the wild relatives, *Beta maritima* best fits these criteria. In its native habitat, *B. maritima* persists in numerous environments. Its adaptation to this range of environments has resulted in the accumulation of stress response traits different from cultivated beet. Over the past 20 years many representatives of this species have been collected, preserved, and made available to breeders. Several breeders (Manerati, Dahlberg, Lewellen, and Doney) have successfully incorporated genes from this wild form into sugarbeet.

The objective of this research project is the development of populations that incorporate some of the genetic diversity from wild *Beta* into sugarbeet. The goal is to produce populations with root characteristics and sucrose concentrations similar to commercial sugarbeet.

Crosses Between Released Fargo Lines and L19

Y317, y318, y322, and y387 are released germplasms (Doney, 1995) derived from the cultivated / maritima cross, L53cms / PI 546420. PI 546420 was collected near Thessaloniki, Greece in 1978. It is a multigerm, non-O type, annual with prostrate growth habit. Testcross hybrids between the released lines and L33 were deficient in sucrose concentration, compared with commercial hybrids. Because of this, it was decided to cross the above germplasm lines to L19 (Theurer, 1978). L19 is noted for its ability to produce hybrids with relatively high sugar concentrations. Its parentage includes the Polish variety 'Udyca'.

Fifty-six families (entries) were grown at Prosper, North Dakota in 1996. Each entry traced back to a single selfed F_1 plant with the pedigree: L53cms / PI 546420 // L19. These families had an average sugar content of 13.3%; ranging from 8.4 to 15.9%. Recoverable sugar per ton of beets ranged from slightly below 100 to 298 lbs. per ton with an average of 237 lbs. per ton.

Individual roots of all entries were sampled for sucrose concentration. The mean of the 842 roots sampled was 14.56%. Entry means of the 56 entries ranged from 10.7 to 17.1% sugar. Selection was based upon both family mean and individual root sucrose within a family. The selected families had means greater than 14.4%. Individual root sucrose concentrations ranged from 7.4 to 19.4% prior to selection. Selected roots ranged from 14.6 to 19.4% with a mean sugar percent of 16.1% or 1.6% higher than the unselected roots. There were 339 roots from 30 entries selected for increase.

Each of the 30 selected entries was maintained as an entity. Eight to 15 roots were selected from each entry for increase in the greenhouse (1997). Seed from plants within an entry (average of 11 plants / entry) was bulked for testing in replicated field tests in 1998. Data from the 1998 trial was of limited value because of conditions related to the extremely wet spring of 1998.

Twenty-four of these 30 families were evaluated in replicated trials again in 1999, using remnant seed from the 1997 greenhouse increase. A number of the lines appear to have higher sugar concentrations than the wild/cultivated parent but are not yet equal to most commercial hybrids (Table 1). Individual roots from 14 of the lines were sampled for sugar concentration. Of the 336 roots sampled 188 were selected for increase and further evaluation in 2000 field trials. Sugar concentrations of the selected roots ranged form 12.2 to 18.5%. All selected roots have acceptable root size and shape. Some beets with sugar concentrations lower than desired were retained to provide a sufficient number of plants for the increase of a line. Depending on the outcome of the 2000 trials, we will either continue selecting within lines or will inter-pollinate lines with sugar concentrations very close to the concentrations observed in standard commercial hybrids.

Crosses of Miscellaneous wild Beta with Sugarbeet

The sugarbeet parent in these crosses was a California line (3747) segregating for genetic male sterility. Crosses were made on male sterile segregates. In subsequent intercrosses, seed was harvested from male sterile segregates to maintain the sterility and insure intercrossing. After two cycles of random intercrossing all populations were grown in a space planted nursery and selected for root shape. Lines that performed well in test crosses (L33cms) in 1996 were increased and evaluated again in replicated trials in 1997. Eleven of the 18 lines tested were increased in the summer of 1998. These were evaluated as lines in replicated trials in 1999 (Table 2). While progress has been achieved in obtaining a more desirable plant and root type, none of the lines have the sugar concentration needed for use in commercial programs. Nine of the eleven lines evaluated in 1999 are being increased in the greenhouse. These will be evaluated again in the field in 2000 and backcrossed to cultivated sugarbeet. F1010 (Campbell, 1990), F1012, or F1013, or F1014 (Campbell, 1992) probably will be used as the sugarbeet parents. These four sugarbeet germplasms have relatively high sugar concentrations and are not derived from the tradition commercial breeding pools.

Recent Introductions to the Breeding Program

Populations were formed by crossing a self incompatible sugarbeet line from California (R376-43) with thirty-seven wild *Beta* accessions from the United Kingdom, France, Ireland, Denmark, Belgium, and the Channel Islands. Ten plants from each wild accession were crossed (as pollinators)

Table 1. Performance of L53/PI 546420//L19 lines at Prosper, North Dakota, 1996 and Fargo, North Dakota, 1999.

		19	96			1999		
		Individual l	Root Sugars				Individual I	Root Sugars
Pedigree	Designation	Before Selection	Selected Roots	Sugar	Root Yield	Recoverable Sugar	Before Selection	Selected For 2000
			%		tons / acre	lbs / acre		%
Y317/L19	C-187**	15.1	16.0	14.0 a-g*	14.0 i-o	3068 fg	13.4	14.6
	C-189**	15.0	15.8	13.7 a-i	14.5 i-m	3181 e-g	13.3	14.3
	C-191**	15.1	15.7	14.2 a-f	12.2 l-o	2982 f-h	12.4	13.6
	C-193**	16.1	16.6	13.7 a-1	10.7 n-q	2253 g-j	14.2	14.8
	C-194**	16.1	16.6	14.6 ab	7.8 q	1709 ij	13.7	14.4
	C-195	14.4	15.3	12.9 g-l	15.5 g-m	3049 fg		
	C-197	17.1	17.2	11.7 lm	13.0 k-o	2249 g-j		
	C-200	16.0	16.1	12.6 i-m	16.6 f-k	3200 e-g	****	
	C-201	15.4	16.0	13.2 d-k	11.8 m-q	2449 g-j	****	
	C-202	15.4	15.8	12.4 j-m	10.4 n-q	2055 h-j		
Mean		15.6	16.1	13.3	12.6	2620	13.4	14.3
Y318 / L19	C-203	15.2	15.5	13.4 b-h	14.0 i-o	2793 f-h	4-	
	C-204**	16.2	16.2	13.7 a-i	14.7 h-m	2950 f-h	13.5	14.0
	C-208**	16.1	16.6	13.7 a-i	12.8 k-o	2654 f-i	13.5	13.9
	C-211**	14.2	15.5	13.7 a-i	13.4 ј-о	2825 f-h	14.1	15.0
Mean		15.4	16.0	13.6	13.7	2806	13.7	14.3
Y322 / L19	C-40	15.1	16.0	12.9 g-k	15.6 g-m	2974 f-h		
	C-45**	15.3	15.9	13.4 b-j	15.6 g-m	3129 e-g	13.7	15.0
	C-62**	15.3	16.0	13.3 с-ј	17.8 f-i	3545 d-f	13.0	13.5
	C-71**	16.2	16.9	13.8 a-i	16.5 f-l	3443 d-f	13.8	15.2
Mean		15.5	16.2	13.4	16.4	3273	13.5	14.6
Y387 / L19	C-76	15.0	15.4	12.0 k-m	19.3 e-g	3634 d- f		
	C-78**	16.8	17.2	13.6 b-j	16.6 f-l	3548 d-f	13.7	15.0
	C-85	14.5	15.1	12.6 h-m	17.4 f-j	3150 e-f		****
	C-89**	15.3	15.5	13.0 g-k	15.2 g-m	2961 f-h	13.0	13.9
	C-92**	15.4	15.8	13.9 a-i	13.7 ј-о	2902 f-h	13.5	14.7
	C-121	15.0	15.9	13.1 e-k	13.9 i-o	2702 f-h		
Mean		15.3	15.8	13.0	16.0	3150	13.4	14.5
Mean all expe	rimental lines	15.5	16.0	13.3	14.3	2892		
Mean lines sel	ected for 2000			13.7	14.0	2939	13.5	14.4
Parents	y317			13.1 g-k	11.9 m-p	2250 g-j		
	y318			11.5 m	9.9 o - q	1642 j		
	y322			13.4 b-j	19.0 e-n	4102 с-е		
	y387			12.8 g-l	17.8 f-i	3616 d- f		
Checks	AC-309		****	14.4 a-d	26.0 bc	5902 a		
	B-3712			14.6 a-c	24.3 b-d	5679 a		
	V-66156			13.5 b-j	26.7 b	5664 ab		
	F1010			13.8 a-i -	20.0 d-f	4228 cd		•

^{*} Means within a column followed by the same letter are not significantly different; LSD 0.10.

^{**} Indicated line was selected for further evaluation or as parental material for future crosses.

Table 2. Yield of "cultivated/wild" sugarbeet, Fargo, North Dakota, 1997 and 1999.

		Su	Sugar	Root Yield	Yield		Recovera	Recoverable Sugar	
Pedigree	Designation	1997	1999	1997	1999	1997	1999	19997	1999
			%	T/A	A	LBS/T-	/ T	——LBS/A	/ A
3747 / B. maritima (Denmark)	C-19*	12.3	10.2	7.1	11.7	210	143	1538	1704
3747 / B. maritima (Belgium)	C-22*	12.3	10.8	7.5	9.5	208	160	1558	1542
	C-153	11.3	9.4	7.8	6.4	185	122	1365	800
3747 / B. maritima (Ireland)	C-27*	12.0	10.7	10.5	10.8	208	156	2200	1700
	C-24*	10.9	11.0	10.5	10.9	180	167	1846	1811
3747 / B. maritima (Middle East)	C-145*	11.2	6.7	11.5	11.7	189	144	2170	1670
3747 / B. Atriplicifolia	C-180*	12.1	10.9	11.4	12.6	205	156	2327	2058
	C-165	11.7	9.4	10.1	6.7	197	131	1933	892
	C-141*	11.0	10.5	11.3	10.7	172	148	1804	1624
3747 / B. macrocarpa	C-29*	10.6	6.7	10.2	11.4	176	136	1789	1583
3747 / B. patula	C-143*	10.9	10.6	12.9	11.1	174	160	2206	1785
F1010		14.2	12.5	9.1	12.0	253	203	2312	2463
VDH - 66140	-	13.7	12.2	15.3	13.9	246	196	3755	2709
ACH-102	1	12.5	11.1	15.7	15.7	210	162	3390	2520
Mean	i	11.5	10.7	10.3	11.6	192	159	1997	1876
LSD (0.10)		1.2	1.0	3.2	3.5	32	26	645	742

* Indicates line was selected for further evaluation or as parental material for future crosses.

individually to R376-43. Ten F_1 plants from each cross (100 plants) were intercrossed to produce the F_2 generation. Equal numbers of seeds from each F_2 plant were grown and intercrossed to produce the F_3 seed. Selection for root shape was initiated with the 1998 crop. Selected plants were increased in the greenhouse to produce seed for a second cycle of selection for root shape in 1999. Plants selected in 1999 will be increased in the greenhouse and subjected to a third cycle of mass selection for desirable plant and root characteristics. Reducing the frequency of plants with multiple crowns may be as difficult as obtaining an acceptable root shape.

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IDENTIFICATION OF SUCROSE METABOLIZING ENZYMES RESPONSIBLE FOR SUCROSE LOSSES DURING SUGARBEET DEVELOPMENT AND STORAGE

BSDF Project 650

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Three enzymes are responsible for nearly all sucrose degradation in sugarbeet. Acid invertase, alkaline invertase and sucrose synthase degrade sucrose to the metabolically active hexose sugars. Invertases catalyze the hydrolysis of sucrose to produce the two invert sugars, glucose and fructose. Invertases are categorized into two groups based on their pH optimum for activity (Tymowska & Kreis, 1998). Acid invertases are most active at pH 4.5 to 5.5, and occur as soluble isoenzymes located in the cell vacuole or insolubilized in the cell wall. Alkaline invertases are most active at pH 7.0 to 8.0 and are located in the cell cytoplasm. The function of invertases is presently unclear, although it has been suggested that acid invertase is detrimental to sucrose accumulation during root development and may be involved in storage losses (Berghall *et al.*; 1997, Wyse, 1974). No functions are known for alkaline invertase. Sucrose synthase is the other major sucrose degrading enzyme in sugarbeet roots. Sucrose synthase catalyzes the cleavage of sucrose using uridine diphosphate (UDP) to form UDP-glucose and fructose in a reversible reaction. Like alkaline invertase, this enzyme is found in the cell cytoplasm. While its function in sugarbeet is unknown, there is evidence from other plant species that sucrose synthase activity is important for sucrose transport and carbohydrate accumulation in storage organs (Zrenner *et al.*, 1995).

Understanding the role of these enzymes in sucrose losses during root development and postharvest storage has proven difficult due to the nature of the enzymes involved. All the major sucrose degrading enzymes exist not as single enzymes, but as families of related isoenzymes. Although isoenzymes are broadly similar in their reactivities, they are typically expressed at different stages of development, have different biochemical properties and are likely to perform different roles in the plant. To better understand the role of the major sucrose degrading enzyme activities in sucrose losses in sugarbeet roots, a study of the activity of individual sucrose degrading isoenzymes was initiated. Specifically, this research has sought to determine the number of isoenzymes of the major sucrose degrading enzymes in sugarbeet roots and their relative contribution to sucrose degradation during root development and postharvest storage.

Methods

Sugarbeet hybrid H66156 (Van der Have) was used in all studies except for the respiration study of field grown sugarbeet roots in which the sugarbeet hybrid 9363 (Maribo) was used. For the developmental study, plants were greenhouse grown with supplemental lighting and 16 hr days. For postharvest study, field grown roots were harvested 120 days after planting, washed and stored at 6, 12 or 21°C and 95% relative humidity. Ten replicate roots were collected for each sample. Soluble proteins were extracted from root samples by homogenization of lyophilized tissue in extraction buffer (100 mM HEPES-NaOH, pH 7.2, 10 mM Na₂SO₃, 5 mM DTT and 1mM MgCl₂) and centrifugation to remove cell debris. Crude extracts were dialyzed overnight against 10 mM

HEPES-NaOH, pH 7.2, 1 mM DTT and 1 mM MgCl₂ to remove sugars. The protein extracts were assayed for acid and alkaline invertase activity by the method of Goldstein and Lampen (1975) at pH 4.7 and 8.0 for acid and alkaline invertase, respectively. Sucrose synthase activity was measured in the direction of sucrose breakdown by the reducing sugar assay of Somogyi (1952). Insoluble acid invertase activity was measured in protein extracts from the cell wall or by direct assay of the cell wall pellet. For extracted cell wall proteins, the pellet of cell debris was washed twice with extraction buffer, extracted overnight with cell wall extraction buffer (100 mM HEPES-NaOH, pH 7.2, 10 mM Na₂SO₃, 5 mM DTT, 2 M NaCl and 15 mM EGTA), centrifuged to remove cell debris and dialyzed overnight. Cell wall invertase activity was assayed as described above for soluble acid invertase.

The presence of isoenzymes for each enzyme family was determined by activity staining of isoelectric focused polyacrylamide gels with ampholines in the pH range of 3.5 to 9.5. Focused gels were incubated for 30 minutes in substrate and stained with 0.1% (w/v) 2,3,5-triphenyltetrazolium chloride (Gabriel and Wang, 1969). Substrates used were 100 mM sucrose for invertase activity and 100 mM sucrose and 10 mM uridine diphosphate for sucrose synthase activity. Acid invertase, alkaline invertase and sucrose synthase activities were assayed at pH 4.7, 7.8 and 6.5, respectively. Control gels were incubated in the appropriate buffer without substrate and stained as above.

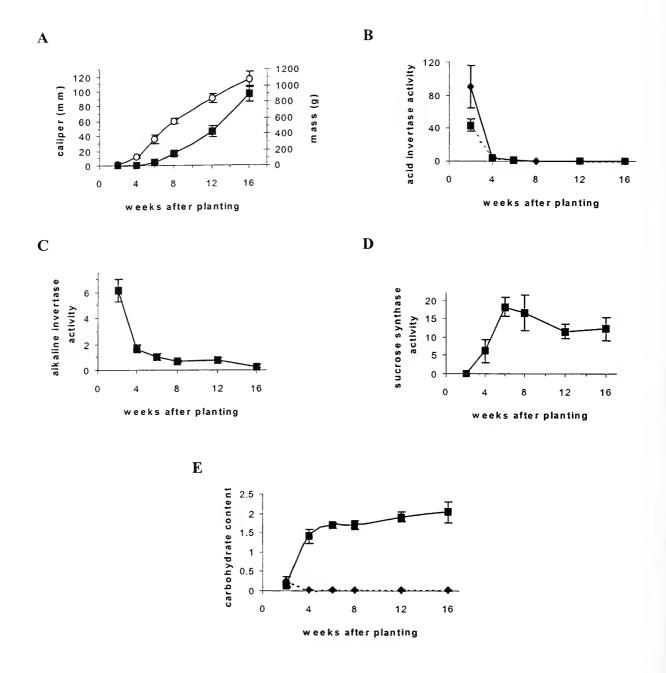
Sucrose, glucose and fructose contents were determined by HPAE-PAD using lactose as an internal standard. Soluble carbohydrates were extracted twice with refluxing 80% EtOH. After evaporation of EtOH, the extract was passed over a bed of C₁₈ and eluted with H₂O. The eluate was filtered through a 0.22 µm nylon filter and injected onto a Dionex CarboPak PA-10 column. Carbohydrates were eluted isocratically with 60 mM NaOH at 1.0 ml/min and detected with an electrochemical detector operating in pulsed amperometric mode.

Respiration was measured by placing six to eight roots of known weight into a sealed six gallon bucket with a continuous air flow of 350 ml min⁻¹. After 24 hr, the CO₂ exiting the bucket was measured using an infrared CO₂ analyzer. Respiration was corrected for background CO₂ levels by measuring the CO₂ exiting an empty bucket and corrected for standard temperature and pressure. Three replicate buckets were measured for each data point.

Results

Developmental Study

The relative contribution of acid invertase, alkaline invertase and sucrose synthase to the total sucrose degrading activity of sugarbeet roots changes with root development. Similarly, the contribution of individual isoenzymes of these enzyme activities to sucrose degradation also changes with development. Figure 1 shows the change in total activity for the three major sucrose degrading activities of sugarbeet roots during development in relation to root size and carbohydrate accumulation. In young roots, the invertases are the predominant sucrose degrading activities. Soluble acid invertase, cell wall acid invertase and alkaline invertase were all at their highest levels in two week old seedlings. The greatest sucrose degrading activity in seedling roots, however, was soluble acid invertase. Only one isoenzyme was responsible for this activity. Its activity was more than double the activity of extractable cell wall acid invertase activity and nearly fifteen times greater



- A. Change in root caliper, measured at widest portion of root $(- \circ -)$ and mass of whole root $(- \blacksquare -)$.
- **B.** Change in soluble acid invertase activity (- -) and extractable cell wall acid invertase (- -).
- C. Change in alkaline invertase activity.
- **D.** Change in sucrose synthase activity.
- E. Change in sucrose content $(-\blacksquare -)$ and reducing sugars $(-\blacksquare -)$. Reducing sugars is combined concentration of glucose and fructose.

Figure 1: Change in root size, acid invertase activity, alkaline invertase activity, sucrose synthase activity and carbohydrate content during sugarbeet root development. Activity for all enzymes is expressed as µmol sucrose mg protein-1 hr-1. Carbohydrates are expressed as mmole g dry wt-1.

than the activity of alkaline invertase. Beyond the two-week stage, invertase activity declined precipitously. Both soluble and extractable cell wall acid invertase activities declined 22- and 12-fold, respectively, between two and four weeks after planting, and by six weeks, their activities were barely detectable. Alkaline invertase activity decreased slightly between two and four weeks and was present at low, relatively constant levels throughout subsequent sugarbeet root development. Two alkaline invertase isoenzymes with isoelectric points of 5.3 and 5.9 contributed to this activity. Although both isoenzymes were present throughout root development, their relative contribution to total activity changed with age. As sugarbeet roots matured, the contribution of the more anionic of these two isoenzymes to total alkaline invertase activity increased, while the activity of the more cationic isoenzyme decreased. Sucrose synthase was the predominant sucrose degrading enzyme during all but the earliest stages of growth and accounted for nearly all sucrolytic activity in mature sugarbeet roots. Sucrose synthase activity increased during the first six weeks of growth and remained at high levels for the remainder of the growing period. Two sucrose synthase isoenzymes contributed to sucrose synthase activity. Only one isoenzyme was evident in roots during the first twelve weeks of growth. Two isoenzymes were present by sixteen weeks.

Postharvest Study

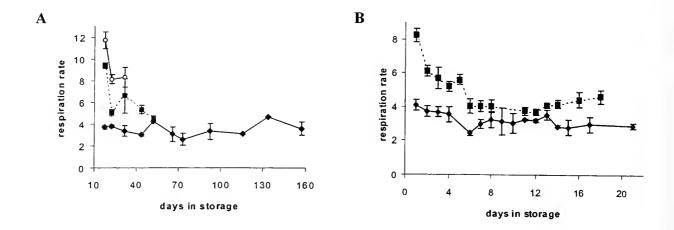
The change in sucrose degrading enzymes during storage under favorable and unfavorable conditions is ongoing. Change in total enzyme activity and isoenzyme levels are being measured in field grown sugarbeet roots stored at 6, 12 and 21° C. Although studies are not complete, initial results suggest only minor changes in total acid invertase, alkaline invertase and sucrose synthase activities during storage. Sucrose synthase remains the major sucrose degrading enzyme throughout storage. Soluble acid invertase, cell wall acid invertase and alkaline invertase are present at low levels even after prolonged storage or storage at elevated temperatures.

The respiration of sugarbeet roots at three different storage temperatures was also measured (Figure 2). Respiration is thought to account for 50 to 70% of sucrose losses in storage (Wyse & Dexter, 1971). Respiration of field grown roots and greenhouse grown roots were measured at 6, 12 and 21° C, and 6 and 13° C, respectively. Respiration rate over time in storage was biphasic. The initial phase, occurring in the first seven to fourteen days after harvest, was characterized by a nearly linear decline in sugarbeet root respiration. The duration of this stage was shorter for greenhouse grown sugarbeet roots (Figure 2B) than for field grown roots (Figure 2A) and probably reflects the gentler harvest and handling conditions these roots received. After the initial period of declining respiration rate, a second phase of respiration was observed during which sugarbeet root respiration remained relatively constant, even after prolonged storage. Root respiration rate during this phase was dependent on storage temperature.

Discussion

Different sucrose degrading enzymes are important at different developmental stages. In young roots, the invertases, especially the acid invertases, are the predominant sucrolytic enzymes. Their contribution to the total sucrose degrading activity in roots, however, is minimal after six weeks of growth. By six weeks, sucrose synthase is the major sucrose degrading activity and remains the major sucrose degrading activity in all subsequent stages of development. It is during this period, when sucrose synthase is most active, that sucrose accumulation in the root is greatest. Sucrose losses during this period, therefore, are most likely to occur by the action of one or more sucrose

synthase isoenzymes. Sucrose synthase also appears to be the major sucrose degrading enzyme during postharvest storage, although these studies are not yet complete. It most certainly is the predominant sucrolytic enzyme in the first two weeks after harvest when root respiration is greatest.



A. Respiration of field grown sugarbeet roots stored at 6°C (— ◆ —), 12°C (— ■ —) and 21°C (— ○ —).

B. Respiration of greenhouse grown sugarbeet roots stored at 6°C (— ◆ —) and 13°C (— ■ —).

Figure 2: Respiration rate (ml CO₂ kg⁻¹ hr⁻¹) of sugarbeet roots stored at different temperatures.

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SUGAR BEET RESEARCH

1999 REPORT

Section E

Sugarbeet and Bean Research Unit Agricultural Research Service - USDA East Lansing, Michigan

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I. SUGAR BEET ACTIVITIES OF THE USDA-ARS E AST LANSING CONDUCTED IN COOPERATION WITH S AGINAW VALLEY BEAN AND BEET FARM DURING 1999.

The USDA-ARS conducted four trials at the Saginaw Valley Bean and Beet Research Farm, Saginaw, MI in 1999. Two of the trails used the same accessions in different locations for seedling disease evaluation (Tests 9911BB and 9913BB, reported together). Two other trials were the standard agronomic test (9912BB, reported here), and a *Cercospora* trial planted alongside of the Michigan and Monitor Sugar Cos. *Cercospora* variety evaluation (not reported here).

The 1999 sugarbeet field trials were planted in Range 9, tiers 7 through 10. This land had been in corn in 1999. The soil was prepared by fall plowing, followed by frost tillage in the early spring. Beets were planted on April 29, 1999. Pre-emergence herbicide (3 qt. Pyramin and 2 qt. Nortron SC per acre) was banded onto the rows immediately following seeding. Seed germination was good overall. Plots were thinned to 6 to 8" between plants within the row and weeded by the second week of July, resulting in good plant stands after thinning and weed control. All experiments were machine harvested October 5, 1999. Sugar analysis was generously provided by the Michigan Sugar Co. sugar laboratory and their assistance is greatly appreciated. All statistical analyses were performed with the aid of MSTAT and / or JMP (SAS Institute). *Cercospora* was controlled with applications of Benylate, Super Tin, and / or Manzate.

TESTS 9911BB AND 9913BB: FIELD EVALUATION OF EMERGENCE UNDER SEEDLING DISEASE PRESSURE

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The objective of this test was to examine field emergence in a range of *Beta* germplasm to evaluate for early resistance to seedling diseases. 114 entries were drawn from germplasm held in the USDA National Germplasm System, and an additional 11 from recent East Lansing releases (Table 1). Generally, the Plant Introduction (PI) accessions were chosen from geographic regions of collection, reasoning was that accessions from warmer and drier areas may show either better tolerance of abiotic stress (including water stress early in the season) or to higher temperature seedling diseases such as *Aphanomyces* and *Rhizoctonia*. Due to limited seed quantities of the PIs, only a single replication of 150 seeds was evaluated in each of two treatments (e.g. 9911BB and 9913BB). Approximately 200 seeds from the seed received are currently available for re-testing. ACH 555 was used as an external check.

Test 9913BB was planted in the south half of Range 7, tiers 11-12. These plots have a history of poor beet growth, presumably due to high seedling disease pressure. All major groups of seedling disease fungi were isolated here in 1999, with *Pythium* being particularly prevalent (John Halloin, pers. comm.).

Emergence counts were taken five times between the time of first emergence on May 12 until June 10. Each plot was planted with 150 seeds as received from the GRIN system. Stands were not thinned, nor was yield data taken. Emergence data was analyzed prior to selections for further evaluation. Selections were based on a number of criteria, including (i) high stand persistence relative to maximal emergence in both tests, (ii) high persistence in 9911BB (e.g.

good ground) and 9913BB (e.g. disease plots), and (iii) final stand evaluations in September 1999. Accessions with reasonable emergence scores but showing severe *Rhizoctonia* damage were excluded from selections at this time, as were accessions that flowered during the season. 23 lines were selected from 9913BB for crossing in the 2000 greenhouse (Table 1).

Results: This test was conducted to determine the feasibility of seedling disease resistance testing by comparing within and between diseased and non-diseased plots at the B&B Farm. As a first approximation, the test was successful for 1999 and it appears that tests like this may serve as a first screening to identify potential germplasm sources to combat seedling diseases. Emergence counts are not presented here, but will be made available on request. Thus, this report presents an overview of the trials.

Excellent stands were obtained in 9911BB (good ground) for all but one accession (PI 558505), but stands were variable in 9913BB (diseased ground) and mortality was higher. Growth throughout the season of all accessions was also superior in 9911BB vs. 9913BB, although precise measurements were not taken for relative growth rates nor biomass accumulation. Some accessions showed greater emergence values in 9913BB relative to 9911BB, but this was likely due to greater moisture availability in 9913BB during the early part of the season since these plots are slightly lower in elevation than 9911BB.

Stand loss was observed in all plots of both 9911BB and 9913BB. In most cases, half of the seedlings counted at maximum emergence were present by the last count in 9911BB, and perhaps slightly fewer in 9913BB. The lack of persistence presented problems with interpreting emergence data, since no clear persisting PIs were evident by the fifth count even in the good ground. By the fifth count, almost without exception, accession counts were lower in the disease plot than the non-disease plot.

All accessions were described as cultivated biennial types in the GRIN system, but 24 of the 125 lines flowered under conditions at the Bean and Beet Farm in 1999. These annual types were rogued as soon as possible after flowering. At least one of the annual types (PI 163176) showed promising emergence results from the analyses. This accession will need to be reevaluated since no plants were brought back for crossing.

In general, Eastern US Germplasm performed among the best as a group. Their stands were more uniform than any other group. ACH 555 showed similar performance. However, in the disease test, plant size was markedly reduced suggesting that the continuous disease pressure was detrimental for not only emergence but also subsequent growth and development.

One exceptional accession was evident in the disease test group, PI 590770, which turned out to be SP85303. SP85303 was developed by G. Coe of the USDA-ARS at Beltsville and is among the most resistant *Aphanomyces* selections developed by him. In the 1992 Bean and Beet Farm Report, SP85303 (reported as 88EL303) had reasonable performance in a standard agronomic test with 16.89% sucrose and 21.8 tons per acre. Although weights were not taken from the 1999 disease plot, the SP85303 beets harvested were superior in size, lack of disease lesions, and relative weight compared with all other selections.

Table 1: Accessions tested for field emergence under seedling disease pressure. Asterisks indicate an accession that behaved as an annual, bold indicates lines selected.

ID	ПЕМ	ORIGIN	TYPE	ID	MEM	ORIGIN	TYPE
Ames 2684	Ames 2684	nd	nd	PI 174063	KOCABAS	Turkey	FODDER
Ames 3062	Ames 3062	Denmark	nd	*PI 175047	PALAK	India	LEAF
Ames 8288	B180	UK	nd	Pl 175594	No. 5973	Turkey	SUGAR
Ames 8289	B182	UK	nd	PI 175597	KOCABAS	Turkey	FODDER
Ames 8294	B197	UK	nd	PI 175598	KOCABAS	Turkey	FODDER
PI 109040	No. T-169	Turkey	FODDER	PI 175599	KOCABAS	Turkey	SUGAR
PI 117116	No. 296	Turkey	FODDER	PI 175600	KARACA OREN	Turkey	FODDER
PI 117117	No. 299	Turkey	SUGAR	PI 175601	PAZI	Turkey	SUGAR
PI 120282		Turkey	FODDER	PI 176423	KOCABAS	Turkey	SUGAR
PI 120689	No. 1219	Turkey	FODDER	PI 176424	PAZI	Turkey	SUGAR
PI 120695	No. 1814	Turkey	FODDER	PI 176425	No. 8972	Turkey	nd
*PI 120696	No. 2124	Turkey	SUGAR	PI 176426	KOCABAS	Turkey	FODDER
PI 120704	No. 3170	Turkey	FODDER	PI 177273	No. 6361	Turkey	FODDER
PI 120705	No. 3208	Turkey	SUGAR	PI 177274	No. 9763	Syria	TABLE
PI 120707	No. 3264	Turkey	FODDER	PI 177275	BELEDI	Turkey	TABLE
PI 124528	CHAKUNDA	India	TABLE	PI 178837	PAZI	Turkey	FODDER
PI 140357	No. 6820	Iran	FODDER	PI 179173	No. 5016	Turkey	SUGAR
*PI 163176	PALOG	India	LEAF	*PI 179179	CICLA	Turkey	FODDER
PI 163178	CHOGHUNDUR	India	TABLE	*PI 181011	No. 8563	India	LEAF
PI 163179	CHOGHUNDUR	India	TABLE	PI 181859	CICLA	Syria	TABLE
PI 163182	CHOGHUNDUR	India	TABLE	PI 181930	Homs No. 30	Syria	TABLE
PI 164292	No. 8928	India	TABLE	PI 181931	CICLA	Syria	LEAF
PI 164659	No. 9084	India	TABLE	PI 204677	No. 174	Turkey	FODDER
*PI 164747	SAG	India	LEAF	PI 204678	No. 178	Turkey	FODDER
PI 164805	CHOGHUNDAR	India	TABLE	*PI 206407	No. 694	Turkey	FODDER
*PI 164806	PALAK	India	LEAF	*PI 212883	PALAK	India	FODDER
PI 164810	No. 9240	India	LEAF	*PI 212884	PALAK	India	LEAF
PI 164968	No. 44	Turkey	TABLE	*PI 215577	No. 13676	India	FODDER
PI 165013	HAYVAN PAU.	Turkey	SUGAR	*PI 217964	RALEK	Pakistan	LEAF
PI 165037	No. 113	Turkey	FODDER	*PI 264150	INDIA	India	LEAF
*PI 165502	PALAK	India	LEAF	*PI 269871	No. 421	Pakistan	LEAF
PI 169014	No. 1394	Turkey	SUGAR	*PI 269872	No. 507	Pakistan	LEAF
PI 169015	No. 1423	Turkey	TABLE	PI 269873	CHINA	Pakistan	SUGAR
PI 169016	PAZI	Turkey	SUGAR	*PI 269874	No. 698	Pakistan	LEAF
PI 169018	PANCAR	Turkey	SUGAR	PI 269875	No. 920	Pakistan	SUGAR
PI 169019	No. 1844	Turkey	TABLE	*PI 271438	PALAK	India	LEAF
PI 169020	PAZI	Turkey	SUGAR	PI 271439	1189	India	TABLE
PI 169023	No. 2246	Turkey	TABLE	*PI 271440	1276	India	LEAF
PI 169024	KIRMIZI	Turkey	SUGAR	*PI 271441	1286	India	LEAF
PI 169025	No. 2693	Turkey	SUGAR	*PI 275637	1349	India	LEAF
Pl 169027	No. 2952	Turkey	FODDER	*PI 277270	BANERJEE'S GIANT	India	LEAF
PI 169028	No. 2960	Turkey	TABLE	PI 285589	EPIPSKI FREEGE	Poland	TABLE

Table 1 (con't): Accessions tested for field emergence under seedling disease pressure.

ID	ITEM	ORIGIN	TYPE	ID	ΠEM	ORIGIN	TYPE
PI 169029	PANCAR	Turkey	TABLE	PI 285590	EPIPSKI HOSER	Poland	TABLE
PI 169030	No. 3395	Turkey	TABLE	PI 285591	OKRAGLY CIEMNOCZ.	Poland	TABLE
PI 171509	HAYVAN	Turkey	FODDER	PI 285592	CRASSA STRZELECKI.	Poland	SUGAR
PI 171512	No. 6864	Turkey	FODDER	PI 285593	CRASSA UDYCKI ZOL.	Poland	SUGAR
PI 171513	No. 6883	Turkey	FODDER	PI 285594	CRASSA WALCOWAT.	Poland	SUGAR
PI 171516	No. 7154	Turkey	FODDER	PI 285595	CRASSA WALCOWAT.	Poland	SUGAR
PI 171517	No. 7159	Turkey	SUGAR	PI 293419	PODZIMNIAJA 0474	F. USSR	TABLE
PI 171518	No. 7164	Turkey	FODDER	PI 357357	OKRUGLA	Macedonia	RED
PI 172730	No. 7425	Turkey	FODDER	PI 357360	Ohridska Zolta	Macedonia	RED
PI 172740	KOCABAS	Turkey	FODDER	PI 357366	Zolta	Macedonia	LEAF
PI 172741	No. 8490	Turkey	LEAF	*PI 408965	Pusa Jyoti	India	nd
PI 173842	CHOGHUNDAR	India	TABLE	PI 546390	WB 69	us	WILD
*PI 173843	PILAK	India	LEAF	PI 546411	Ames 4218	UK	WILD
PI 173844	CHOGHUNDAR	India	TABLE	PI 558505	FC 506	us	SUGAR
PI 174058	No. 7764	Turkey	FODDER	PI 558515	FC 403	us	SUGAR
EASTERN US GERMPLASM							
PI 590770	SP85303-0	us	SUGAR		SR80	us	SUGAR
	98EL02	us	SUGAR		SR87	us	SUGAR
	98EL04	us	SUGAR		SR93	us	SUGAR
	EL38	US	SUGAR		SR94	us	SUGAR
	EL48	US	SUGAR		SR95	us	SUGAR
	EL50	US	SUGAR		EL51	us	SUGAR
				l			

Figure 1: Relative performance of accessions between plots.

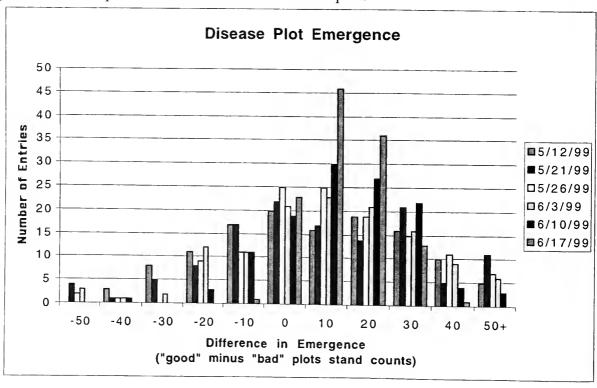


Figure 1 presents the relative performance of accessions in both the "good" non-diseased plots (9911BB) and the "bad" disease plots (9913BB) for each of the six counting dates. Each plot had the same number of seeds planted. Most accessions performed better in the non-diseased plots (e.g. a difference in emergence > 0). Accessions with scores <0 were seen, but their significance is unclear at this point. Of particular interest are the accessions with little or no differences between plots, as these are fairly numerous in number and may present sources of genes that appear to perform with less environmental dependence, including disease pressure, than others.

In total, we were encouraged by the results form comparing emergence and persistence in diseased and non-diseased plots. However, the interpretation of these results must be qualified. First, direct comparison is not possible due to differences in the speed of emergence (or other developmental processes), likely due to available moisture supply. Second, screening in the disease nursery identified accessions that performed better or worse under disease conditions, but the problem of stand persistence in good ground appears <u>not</u> to have been addressed by these comparisons. Third, because PIs differ markedly in the genetic structure of their populations, it is possible that effective and desirable seedling resistance genes are present at low frequency among the survivors. It is apparent that at least some of these genes are present at high frequency in accessions from the Eastern US germplasm pool. And finally, perhaps the most apparent operational criteria for selection is equivalent performance in diseased and non-diseased plots. In general, those accessions that we selected showed more similarities in performance between good and disease plots, suggesting that their performance might be expected to be more consistent across a range of environments.

EXPERIMENT 9912BB: AGRONOMIC EVALUATION OF SMOOTH ROOT RELEASES AND PROSPECTIVE RELEASES – 1999

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This experiment was designed to evaluate performance of 24 entries for the standard agronomic parameters, and smoothroot score as a surrogate measure for low soil tare. We considered this test reliable and well executed and in line with past performances of those entries tested in prior years. Three commercial hybrid varieties (ACH 185, Betaseed 5931, Novartis E17) and two popular hybrids from the 1970s and 1980s (US H20 and US H23) provided performance references.

Three East Lansing releases were included; a West Coast Beet Seed increase of the 1971 release monogerm O-type EL38, 1997 smoothroot release SR94, and 1998 smoothroot release SR95. Five were of the planned smoothroot releases SR96, 94HS25, monogerm multiple disease resistant 99J19-00, monogerm 99J31-00 and monogerm 99J33-00. Two others were 98EL02 and 98EL04 scheduled for release combining parentage and selection for smoothroot, higher sucrose, and the Holly monogenic Rhizomania resistance. Three lines (98J24-01, 98J34-01, and 98J41-01) have common background from hybridizing high sucrose, smoothroot germplasm with Hogaboam era monogerm germplasm containing significant levels of Rhizoctonia resistance plus high levels of Cercospora and Aphanomyces resistance. The list of entries is rounded out by five lines derived entirely or at least 50% from population 95H07, itself a cross of an EL50 root with a selected smooth root beet. Prospective release 99J19-00 is also derived predominantly from 95H07. Also included in this test was an experimental smooth root line from a seed company (97-060515-01).

For ease of reference, these groupings of entries are listed below:

- 5 are released hybrids,
- 1 is an experimental hybrid,
- 3 are past EL releases,
- 5 are prospective smoothroot EL releases,
- 2 are planned smoothroot EL releases with Rhizomania resistance,
- 3 are lines with high sucrose smoothroot and Hogaboam era Rhizoctonia resistance ancestry,
- 5 are 95H07 derivatives (6 including 99J19-00).

A more condensed grouping helpful in interpreting performances is:

- 4 are modern hybrids,
- 7 are entries with traditional rough exteriors,
- 17 are smoothroot entries,
- 3 are entries without Theurer era high sucrose percent ancestry,
- 4 are past or planned SR releases with moderately high sucrose percent,
- 5 are monogerm lines of three different ancestries.

Performance is ranked by recoverable white sugar per acre (RWSA) in Table 1. A conspicuous grouping in the top five entries for RWSA is seen with the East Lansing germplasms SR95, SR96, and SR94, with the remaining two of the top five modern hybrids. Interestingly, SR95,

SR96, and SR94 (RWSA mean of 7312 lbs. / A) each have a diverse ancestry. All four modern hybrids (RWSA mean of 6826 lbs. / A) are ranked in the top half of the entries. The lowest RWSA was O-type EL38, a 1971 Hogaboam self-sterile release likely based on several clones. Next lowest in ranking is 98J24-01, which appears to be at least an S₂ generation family, derived from an unrecognized self-fertile beet selected for smoothroot and high sucrose percent by Clair Theurer at East Lansing (prior to his retirement).

Beet yield by tonnage per acre is not ordered in Table 1, but some patterns are evident. Ranks 1 and 6 of the 24 in the test are held by derivatives of population 95H07. The top rank is held by 98J02x05, a pair cross of two smoothroot selections from 95H07, and the #6 rank is held by 99J19-00, a close relative of 98J02x05. 99J19-00 is closely derived from 98J19-01, which topped the tonnage per acre ranking among fifteen entries in a test at the Saginaw Valley Farm in 1998. (Overall, in the two 1998 tests at the Saginaw Valley Farm, 3 of the top 5 entries for tonnage were 95H07 derivatives.) EL38 is ranked least in tonnage, being somewhat of an inbred. The next to last tonnage ranking belongs to the top sucrose entry 98J24-01, discussed above as an inbred line and seen in the field as a smaller canopy entry.

Sucrose percentage by entry ranged from 18.23 to 15.41, with the three modern commercial hybrid cultivar checks averaging 18.00%. Top ranking was held by the moderately smoothroot 98J24-01 with 18.23%. The spread of 2.8 % points between the lowest entries and the modern commercial checks is similar to that from most other years with full season growth. One grouping of entries by sucrose percent consistent with past years is the "traditional East Lansing germplasm" trio of 98J02x05, 99J19-00, and 99J02-00 with a mean of 15.52 (range was 15.41-15.64). Other East Lansing breeding lines and releases with various proportions of high sucrose percent parentage had a continuum of sucrose concentrations above 16%.

Clear juice purity (CJP%) rankings were topped by modern commercial hybrid Novartis E17 (93.88%), with three (mean = 93.67%) of the four top spots held by three of the four modern hybrids. The six lowest purities were held by the group of various 95H07 derivatives (mean = 91.92%), a pattern also seen with that germplasm in 1998.

Amino N rankings had groupings including both members of the closely related pair (mean = 9.80) of monogerm smoothroot lines 99J31-00 and 99J33-00, ranked 2nd and 4th best, respectively. Overall, test range was 9.26 - 17.57. Mean of the three modern commercial checks was 10.57 (range was 9.26 - 11.82). The six members of the 95H07 derivative group ranked in the worst third of the rankings. Amino N of EL38 ranked 21st, but in retrospect, this may have been due to its poor stand and intrinsic low vigor. Adjustments to tonnage per acre can be figured from pre-harvest stand and gap measurements, but these same adjustments can't be (easily) used to adjust amino N. Poor stand differentially makes more nitrogen available to the beets that are there, delaying the idealized late season transition from nitrogen luxury to nitrogen paucity.

Smoothroot (SR) score rankings showed seven "traditional" sugarbeet entries (the four modern hybrids, plus US H20, US H23, and EL38) bunched at the highest values (i.e. the deepest sutures). Mean of the seven was 2.17, and the range 2.04 - 2.25. The smoothest entries scored 1.50 - 1.60, including SR95, SR96, and prospective releases 94HS25, 99J19-00 and 99J31-00. SR95 was considered the smoothest line entered in the test, from prior years' scores.

The properties of the nine breeding germplasm entries, agronomically evaluated for the first time here, indicate the recent emphasis of the sugarbeet breeding program at East Lansing. All nine

new entries are high to moderate for smoothroot as well as for resistance to *Cercospora* and *Aphanomyces* Four of the nine are monogerm with enhanced CMS-maintainer frequency, moderate *Rhizoctonia* resistance, and/or improved sucrose percent, depending on the entry. The other five new entries carry deliberately introduced recessive alleles for monogerm or CMS-maintenance that can be recovered in fixed form in current or future generations. Emphasis on disease resistance and higher sucrose percentage will continue and complement the easily selectable smoothroot characteristic.

Table 1: Agronomic performance of lines in Test 9912BB.

Entry	RWSA	RWST	Tons/Acre	Sucrose%	CJP %	Amino N	SR score
SR95	7465.6	240.8	31.03	16.98	93.28	12.56	1.54
97-060515-01	7407.0	245.9	30.12	17.25	93.44	9.72	2.21
SR96	7330.6	247.6	29.60	17.53	93.01	13.59	1.50
Betaseed 5931	7201.5	260.1	27.68	18.09	93.68	10.62	2.17
SR94	7142.6	243.4	29.36	17.12	93.37	11.26	1.75
98J34-01	7056.8	234.9	30.03	16.95	92.26	15.41	1.88
98EL04	6961.1	223.3	31.20	16.09	92.51	12.68	1.71
98J02X05	6916.2	212.4	32.62	15.64	91.70	14.23	1.58
98J27-00	6428.2	225.9	28.54	16.41	92.08	16.48	1.63
Novartis E17	6420.9	255.6	25.10	17.73	93.88	9.26	2.21
98J41-01	6329.7	246.1	25.71	17.45	92.94	10.70	1.75
ACH 185	6277.0	254.1	24.67	18.18	92.44	11.82	2.21
99J19-00	6243.4	210.0	29.73	15.51	91.60	17.57	1.50
94HS25	6167.3	240.6	25.59	17.21	92.64	12.30	1.54
99J02-00	6104.5	210.8	28.99	15.41	92.08	14.64	1.75
98EL02	6041.3	226.8	26.49	16.08	93.21	12.85	1.75
97J27-00	6009.9	226.8	26.51	16.51	91.99	13.87	1.50
US H23	5999.1	227.1	26.39	16.46	92.17	13.28	2.13
98J28-02	5867.8	223.5	26.20	16.25	92.08	14.19	1.83
99J33-00	5842.6	224.7	26.02	16.00	93.04	9.88	1.50
99J31-00	5793.8	233.1	24.82	16.51	93.17	9.72	1.63
US H20	5394.8	238.4	22.61	16.73	93.53	11.51	2.04
98J24-01	5091.3	259.1	19.64	18.23	93.15	10.40	1.79
EL38	4069.1	225.7	17.98	16.26	92.47	14.95	2.25
Mean	6315.1	234.9	26.94	16.77	92.74	12.65	1.81
CV	16.43	6.83	16.42	5.20	1.04	26.51	19.13
LSD (0.05)	1584.0	16.21	6.46	0.77	1.71	5.91	0.53

II. Germination of Sugar Beet (Beta vulgaris) Under Stress Environments : A Survey of Differential Gene Expression in vitro

BSDF Project 741

Benildo de los Reyes – USDA-ARS Research Associate and J. Mitchell McGrath

Background

Germination and seedling emergence are fundamental processes that determine potential harvest on sugar beet crops. Poor germination and emergence, due to biotic and abiotic stresses, are major problems with serious economic impact to the sugar beet industry. While external influences of the environment are well documented, and can be managed to a degree, the intrinsic responses of the plant to external signals such as those imposed stress are not well understood. We are particularly interested in these intrinsic responses because they provide the best evidence for the involvement of genes in stress response, and can give information on the identity of those genes and the conditions under which they are expressed. Ultimately, knowing which genes are expressed, and then deducing and proving which genes control or most influence the appropriate response(s) will allow their directed selection, genetic manipulation, and biotechnological utility for improving germplasm performance.

Genetic causes of emergence and stand reduction failures are not very well understood. Previous observations from both laboratory and field experiments (McGrath, BSDF Project 741) suggested the importance of genetics in the expression of seed vigor in the early stages of sugar beet growth. Expression of seedling vigor is influenced by a number of extrinsic and intrinsic components (Kneebone, 1976). Among the major factors that determine the extrinsic component include the seed production and post-harvest environments. These components account for the variability in germination ability between seed lots of the same cultivar and do not reflect the actual vigor potential of the cultivar. Intrinsic components are determined primarily by the genetic make-up of the seed, and likely some effects imposed by the maternal physiology. Our results to date indicate the problem of poor germination and emergence in sugar beet fields are largely abiotic. Stand reductions after maximal emergence, the stand persistence problem, are largely due to disease. These phases are not mutually exclusive, and likely overlap to an undetermined extent.

The inability of certain cultivars and seedlots to adapt to sub-optimal conditions in the germination environment is clear, but responses are difficult to dissect in field grown materials. Previous laboratory experiments involving artificial stress showed significant differences in the ability of sugar beet cultivars to germinate in aqueous solutions supplemented with different solutes. Some adjuvants promote (i.e. 0.3% hydrogen peroxide) or inhibit (e.g. pure water, 350mM NaCl, 200mM mannitol) germination relative to 'traditional' germination on moist filter paper. Among the cultivars studied, USH20 exhibited superior germination under both artificial stress (laboratory) and actual field conditions, compared to two other cultivars (HME17 and ACH185) that exhibited good and poor vigor, respectively. Based on these findings, the major stress factors that significantly affect the expression of sugar beet seed vigor were identified to include the extremes of moisture (flooding and drought), salinity and anoxia (anaerobic stress).

The objective of this research project is to dissect the molecular components determining the expression of seed vigor in sugar beet, through the discovery of cultivar specific, differential gene action in response to sub-optimal germination environments. The isolation, identification and characterization of specific genes that contribute to abiotic stress tolerance expressed during germination is the primary approach. These genes include those that are either induced or repressed in the cultivar USH20 under specific stress conditions. Future work will include comparisons among good and poor stress emergers.

Discovery of genes with potential roles in stress-tolerance at the germination stage will result in better understanding of the physiological and biochemical processes that limit the expression of seed vigor in sugar beet. This will be important in developing strategies to improve seed vigor by genetic engineering. Along with other genes expressed in germinating seeds, the putative stress-related genes are currently being used as markers (restriction fragment length polymorphism or RFLP) to develop a genetic map of the sugar beet chromosomes. This map will not only provide better understanding of the genetic architecture of the sugar beet genome but will also serve as a more direct route to investigate the chromosomal distribution of gene loci with potential roles in the expression of seed vigor.

Experimental Approach

The cultivar USH20, a good stress-germinator, is our model system to identify genes with potential roles in stress-tolerance at the germination stage. The basic strategy has been to compare gene expression profiles under different laboratory environments. Three types of treatments include germination in moist filter paper (standard or control), submerged germination in pure water or solutions containing 350mM NaCl or 200mM mannitol (stressed or negative treatment), and germination submerged in 0.3 % hydrogen peroxide (positive treatment).

Total RNA samples were isolated by guanidine hydrochloride method from seedlings germinated for 4 days. Differential gene expression analyses were done by comparing mRNA (messenger RNA) fingerprints generated by the differential display-reverse transcription polymerase chain reaction (DDRT-PCR) technique (Liang and Pardee, 1992), using the Delta Differential Display kit (Clontech, Palo Alto CA). Differentially expressed (up- and downregulated) cDNA copies (complementary DNA, reverse copied from mRNA) were identified by comparing mRNA fingerprints' relative band intensities in autoradiograms. Candidate cDNAs were cloned in pT-Adv plasmid vector (Clontech, Palo Alto CA) and the inserts were sequenced by dideoxy-termination method in the Long ReadIR4200 automated DNA sequencer (Li-Cor, Lincoln NE). The putative identity of individual stress-induced cDNA was determined by alignment of nucleotide sequences with known genes in genome databases (GenBank, EMBL, DDBJ) using the blast search algortihms (BlastN, BlastX) (Altschul et al., 1997). The expression of the candidate genes was confirmed through northern blot analysis, by probing mRNA (2 ug) isolated from individual treatment with the radiolabeled cloned cDNAs.

Results

Our initial survey of differential gene expression in USH20 resulted in 807 cDNA fragments, which were amplified using 50 combinations of anchored and arbitrary primers (Clontech, Palo

Alto CA). The patterns observed in the mRNA profiles indicated significant changes in gene expression under optimal and sub-optimal environments. Of the total 807 bands observed, 95% corresponded to genes that were expressed in all treatments. The other 5% showed induced expression in response to at least one treatment. Some of these have been cloned and their nucleotide sequences have been obtained. A partial list of these differentially expressed cDNAs and their induction and expression is given in Table 1.

Table 1. Partial list of up-regulated cDNA clones isolated by differential display analysis of germinating sugar beet (cv. USH20). This list includes only the clones whose expressions were confirmed by northern blot analysis.

CloneID	Primer pair	Partial cDNA size	Induction	Putative identity
L1	P1/P1	827 bp	NaCl, H ₂ O ₂	ATPase
L2	P1/P1	668 bp	NaCl, H ₂ O ₂	unknown¹
L5	P5/P5	484 bp	solution	unknown¹
L7	P8/P8	670 bp	solution	unknown
L8	P9/P9	768 bp	$\mathrm{H_2O_2}$	sugar-PO ₄ translocator
L9	T1/P9	625 bp	H_2O_2	cytochrome b
L16	P8/P8	598 bp	solution	unknown
L18	P2/P3	622 bp	solution	pectinacetyl esterase
L19	P2/P4	327 bp	NaCl	unknown
L20	P2/P5	520 bp	filter paper	unknown
L21	P2/P5	322 bp	NaCl	unknown
L22	P2/P5	526 bp	NaCl	unknown
L24	P2/P6	283 bp	mannitol	unknown
L13	T2/P4	420 bp	all	40S ribosomal protein

¹ similarity to unknown protein in Arabidopsis

Where the function of these clones has been surmised from matches in the world genome databases, further discussion is included below, particularly as it relates to potential clues for stress germination mechanisms and targets for further analyses and manipulations.

Sugar-phosphate translocator protein and Cytochrome b

About 0.4% of the total cDNAs were upregulated in response to germination in hydrogen peroxide solution. The identities of two of these genes were positively identified as cDNAs encoding a sugar-phosphate translocator protein (L8) and cytochrome b (L9), based on significant sequence homology of the partial cDNAs with other known genes in genome databases. Both of the genes exhibited high levels of expression in seedlings germinated in 0.3% hydrogen peroxide relative to the other treatments. The patterns of expression were identical in the differential display profile and northern blots (Figure 1). These results were interpreted as a possible indication of coordinate regulation of expression of these two genes and may be related

to a common path of metabolic adjustment when germination occurs in the presence of hydrogen peroxide. Our initial hypothesis is that hydrogen peroxide relieves anaerobic stress under condition of excess water during germination in solution. We propose the following hypotheses to explain the possible physiological relevance of the current results in relation to our initial hypothesis regarding the role of hydrogen peroxide as the source of oxygen during germination.

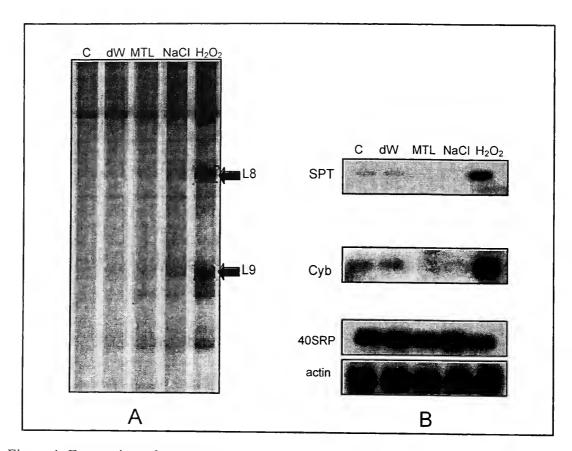


Figure 1. Expression of genes encoding sugar-phosphate translocator protein, SPT (L8) and cytochrome b, cyb (L9) during germination of USH20 in hydrogen peroxide. (A) Differential display of mRNA from seedlings germinated on moist filter paper (control, C), distilled water (dW), 200mM mannitol (MTL), 350mM sodium chloride (NaCl) and 0.3% hydrogen peroxide (H₂O₂). Induced gene expressions are shown by bands L8 and L9 which are present in samples germinated in H₂O₂ but not in the control and other treatments. (B) The bands L8 and L9 were isolated and used as probes to confirm differential gene expression by northern blot analysis. Expression patterns show significant increases in transcript levels corresponding to SPT and cyb genes in the H₂O₂ treated sample. Uniform transcript levels in all samples show constitutive pattern of expression of "housekeeping" genes (40S ribosomal protein/40SRP and actin).

Respiration and oxidative phosphorylation are two of the major metabolic processes that are immediately activated upon imbibition. These processes occur in the presence of oxygen and

provide the ATP (energy) that fuels cellular processes related to germination including cell wall elongation and extension/emergence of the radicle (Nykiforuk and Johnson-Flanagan, 1998)). The increased expression of the cytochrome b gene/s suggests a rapid development of the mitochondrial electron transport system more likely as a consequence of more stable supply of oxygen to the germinating embryo, in the presence of hydrogen peroxide than in solution of pure water. This observation further supports the results from last year's experiments, which indicated that anaerobic stress appears to be a major factor that affects sugar beet germination and emergence.

The substrates for oxidative phosphorylation come from the by-products of sugar metabolism via glycolysis and TCA cycle (Heydecker, 1977). The availability of steady supply of oxygen in the germination solution (hydrogen peroxide treatment) possibly resulted in increased demand for respiratory substrates (feedback control). Based on the current results, we hypothesized that the upregulation of expression of a gene encoding a sugar-phosphate translocator protein is probably related to a mechanism by which respiratory substrates are mobilized to the cytoplasm for glycolysis. Biochemical studies in other plant species indicated the presence of both triose-and hexose-phosphate translocator proteins in amyloplast of non-photosynthetic organs such as the roots and germinating seeds. Unlike their chloroplast counterparts, these proteins transport not only triose-phosphates but also residual hexose-phosphates across the amyloplast envelop to the cytoplasm where they can be utilized for glycolysis (Echeverria et al., 1988; Borchert et al., 1989; MacDonald and Rees, 1983). The possible occurrence of this process in sugar beet (as suggested by the current results), probably ensures that sufficient supply of respiratory substrates are available to meet the energy demand for germination under ideal conditions, an indirect but positive effect of hydrogen peroxide.

ATPase

Salt stress (350mM NaCl) during sugar beet germination induced the expression of several genes. One of the salt-induced cDNAs (L1) was positively identified as that encoding for a membrane-bound ATPase, based on significant sequence homology of the partial cDNA with other ATPase genes in the genome databases. ATPase is an integral component of proton pumps located in the plasma membrane and tonoplast. This protein is involved in active transport of ions from the cytoplasm to the intercellular space and vacuole. Both the differential display profile and northern blot indicated that a putative ATPase gene was highly upregulated during germination in salt and hydrogen peroxide solutions in addition to the basal expression levels in the control (moist filter paper) and other stress treatments (Figure 2). This result suggests that regulation of this gene may be important not only for growth-related processes but also for some stress-related response (Lehr et al., 1999). The potential physiological significance of the salt-induced expression of ATPase gene/s during sugar beet germination may be related to an energy-requiring mechanism that maintains low level of Na⁺ inside the cytoplasm, which could otherwise produce damaging effects to the germinating embryo (Apse et al., 1999; Frommer et al., 1999).

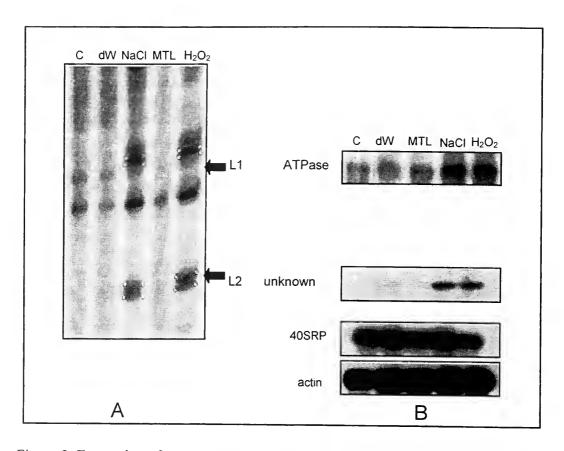


Figure 2. Expression of genes encoding ATPase (L1) and an unknown protein (L2) during germination of USH20 in sodium chloride and hydrogen peroxide.

(A) Differential display of mRNA from seedlings germinated on moist filter paper (control, C), distilled water (dW), 200mM mannitol (MTL), 350mM sodium chloride (NaCl) and 0.3% hydrogen peroxide (H2O2). Induced gene expressions are shown by bands L1 and L2 which are present in samples germinated in NaCl and H2O2 but not in the control and other treatments. (B) The bands L1 and L2 were isolated and used as probes to confirm differential gene expression by northern blot analysis. Expression patterns show significant increases in transcript levels corresponding to ATPase and the unknown gene in the NaCl and H2O2 treated samples. Uniform transcript levels in all samples show constitutive pattern of expression of "housekeeping" genes (40S ribosomal protein/40SRP and actin).

On-going Experiments

The preliminary results from this study suggest that many genes with basic "housekeeping" functions are regulated under stress conditions of germination. The expression patterns of these genes under optimal and sub-optimal germination environments are currently being compared between USH20 (good stress-emerger) and ACH185 (poor stress-emerger) to confirm their direct involvement in cultivar differences in seed vigor.

The putative identities for many of the cDNAs that we isolated are still unknown because of the lack of significant sequence similarity with other genes. These cDNAs are quite interesting because of the possibility that they represent novel genes with important roles in stress-tolerance during sugar beet germination. Recently, we constructed a cDNA expression library (in lambda UniZap-XR vector, Stratagene, La Jolla CA) from pooled mRNA from all six treatments on USH20. This library is currently being screened to isolate the full-length clones corresponding to all the cDNAs listed in Table 1. The full-length coding sequences of these cDNAs will be analyzed for potential structural motifs (at the nucleotide and amino acid sequence levels) that may provide clues regarding the biological role/s of these genes in sugar beet germination and emergence under stress environments. Furthermore, additional genes will be isolated and these efforts will be targeted towards identification of stress-specific genes.

Despite these efforts to isolate and identify genes associated with stress-tolerance during sugar beet germination, we still know very little about the genetic components that determine the expression of seed vigor. However, given the physiological complexity of tolerance to different stresses it is highly possible that hundreds of genes are involved, each one probably associated with specific adaptive mechanism. The result of the current survey of differential gene expression is still far from being comprehensive and does not provide adequate information to better understand the relationship between abiotic stress-tolerance and seed vigor in sugar beet. The initial strategy of using differential display provided useful preliminary information but the results were obviously quite limited in scope. Apparently, a larger scale gene discovery and characterization program will be necessary to satisfy the original goals that were set at the beginning of this research project and to realize the full potential of this project to generate innovative tools for sugar beet improvement.

Our survey of differential gene expression using differential display is still on- going. Additionally, in order to approach this problem in a more functional and global perspective, we are currently developing a small collection of Expressed Sequence Tags (ESTs) from the cDNA library developed from germinating sugar beet seeds. Basically, this collection will be a subsample of partial sequences of the genes that are active during germination under optimal and sub-optimal environments. Our strategy for the generation of this EST collection involves the preferential elimination of the clones from the library that correspond to genes that are not involved with stress- and hydrogen-peroxide induced responses using the methods known as subtractive hybridization and differential screening (Nguyen et al., 1995; Hoog, 1991). The resulting sub-libraries (one each for stress and hydrogen peroxide) represents a snapshot of induced gene expression that will then be characterized by partial sequencing (300-450 bp) from the 5' ends of the individual cDNAs. During the past decade, voluminous amount of gene sequence information had become available in public databases. These databases consist of gene sequence information from both prokaryotic and eukaryotic organisms and include a number of disease causing microorganisms (Saier, 1998), some model plant species like rice (Sasaki et al., 1994) and Arabidopsis (Cooke et al., 1996) and also human (Hillier et al., 1996). These existing databases will be searched for potential homologies with the individual ESTs generated from the subtracted germination-specific cDNA library of sugar beet. This approach will allow a more rapid and direct access to hundreds (or even thousands) of genes and will serve as the foundation for studying global changes in gene expression patterns associated with the ability of sugar beet to germinate under stress. Homologies with other genes or gene motifs whose function had been previously identified will provide wider windows on the molecular genetic basis of seed vigor in a more functional perspective. This approach also offers exciting opportunities to discover novel

metabolic pathways relevant to germination. Future major benefits from this initiative will include opportunities to investigate this problem using more sophisticated tools of genomics including the use of DNA chips and transcription profiling (Roberts et al., 2000). These technologies are predicted to be of huge impact not only to basic research in biology but also to plant breeding and cultivar improvement in the new millennium. Lastly, the small scale EST collection from this project is being used as a foundation to study the genome architecture and evolution of *Beta vulgaris* through the construction of an EST marker-based genetic map of the sugar beet chromosomes.

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GROWTH OF SUGARBEET PATHOGENS IN VITRO.

Joseph W. Saunders

RHIZOCTONIA AND PYTHIUM. The sensitivity of in vitro growth of sugarbeet fungal pathogens *Rhizoctonia solani* (RZT) and *Pythium ultimum* (PYT) to three herbicides (Roundup, Liberty and Pursuit), each affecting separate individual amino acid biosynthetic steps, was evaluated under conditions of both nutritional dependance and independence on inorganic nitrogen from the culture medium. One isolate each of RZT and PYT was grown on various concentrations (0, 2.1-21000 μ M active ingredient) of each of the three herbicide formulations in agar plates with a Murashige-Skoog nutrient medium background, with each of four medium nitrogen regimes: no nitrogen, or nitrogen provided at 30 mM as either casein hydrolysate, ammonium, or nitrate.

For all three herbicides with PYT, nitrogen regime did not affect sensitivity of extension growth of the fungus to the herbicide; PYT was increasingly sensitive to Pursuit, Round-up, and Liberty, in that order. RZT showed the same order of sensitivity to the three herbicides. Nitrogen source only had a significant effect on RZT sensitivity to the herbicide in the case of Liberty, where extension growth on casein hydrolysate as nitrogen source was about tenfold less sensitive to the herbicide than with the three other nitrogen sources. The most noteworthy finding of the entire test was that RZT extension growth at the highest Pursuit concentration (ie, 21,000 μ M), for each of the nitrogen sources, was at least 50% of the growth in the absence of the herbicide. This appears to be a remarkable tolerance of RZT to the herbicide.

This research was prompted by the question of whether RZT or PYT, each a facultative saprophyte, would be sensitive ("fungicidally") to the presence of a herbicide such as might be encountered in field soil, and might thus be controlled to some degree incidentally by herbicides used as part of cell-selection or transgenic herbicide-resistant variety packages. Of the three herbicides, only Pursuit is known to be persistent in soil. However, the absence of lower sensitivity to the herbicide of extension growth with casein hydrolysate (essentially a mixture of amino acids) as nitrogen source in five out of six combinations suggests that herbicide-induced deficiency of one or more amino acids is not an obvious explanation for growth inhibition by the herbicides.

CERCOSPORA. In continued research with Cercospora beticola (CER) inoculated onto Murashige-Skoog plant tissue culture medium, when six CER mycelium plugs were placed on one side of a plate with 1.0 mg/L N⁵-benzyladenine medium, with a leaf disc placed on the other side 13 days earlier, and grown on the lab bench under ambient lighting from ceiling fluorescent lamps, CER growth for two weeks was limited in extent, with no cercosporin production, based on lack of red color in the agar. When CER growth on the surface of the agar ceased, sparse hyphae extended to the leaf disc. Thirteen weeks after inoculation with CER, mycelium had grown on the leaf disc, and produced conspicuous red coloration (cercosporin) in the medium surrounding the leaf disc, but still not in proximity to the dense mycelial masses around the inoculum plugs.

The differential accumulation of cercosporin on different sides of the Petri dish was consistent with a conjectured late presence of organic N or late absence of nitrate from the vicinity of the leaf disc, consistent with cercosporin accumulation we have seen and measured by HPLC from CER on water agar, nitrogen-free Murashige-Skoog medium, and potato dextrose agar medium. Using both defined and undefined complex media, Ehrenshaft and Upchurch

(1993) had reported that host protein(s) induce, and nitrate represses, accumulation of cercosporin in a phytopathogenic strain of Cercospora kikuchii, a pathogen of soybean. This induction of cercosporin accumulation in the vicinity of the sugarbeet leaf disc should be pursued further with a range of germplasm including CER resistance and susceptibility. Differential production of cercosporin by the pathogen in leaf tissue of resistant vs susceptible genotypes could be one mechanism of genetic resistance by the host. Perhaps it also could explain the 'resistance' of young leaves on the host plant.

PUBLICATIONS

J.W. Saunders¹ and C.J. Tsai. 1999. Production of somatic embryos and shoots from sugarbeet callus: Effects of abscisic acid, other growth regulators, nitrogen source, sucrose concentration and genotype. In Vitro Cell. Dev. Biol. –Plant 35:18-24.

Two sugarbeet (Beta vulgaris L.) genotypes, REL-1 and REL-2, were used to measure the level of somatic embryo and shoot production from hormone-autonomous callus plated under varied nutrient medium combinations of abscisic acid with the growth regulators 6benzyladenine, 1-naphthaleneacetic acid, or 2,4-dichlorophenoxyacetic acid, with eight sole nitrogen sources, or with different sucrose concentrations. Clone REL-2 produced embryos up to thirty-five fold more frequently than clone REL-1. Inclusion of abscisic acid at some concentrations consistently improved embryo production in all experiments, and was observed to stimulate shoot production. At some concentrations, 1-naphthaleneacetic acid as well as urea and glutamine stimulated greater embryo production over the control, but only for REL-1, where there was greater room for improvement. Three and five percent sucrose were superior to one, seven, and nine percent. Higher initial 6-benzyladenine concentration (in the range 0, 0.1 - 1.0 mg/L) was associated with lower embryo production but greater shoot regeneration for both clones. REL-2 was significantly better than REL-1 in shoot regeneration. The range of embryo production was more than thirty-five fold between genotypes, whereas the range of physiological effects was no greater than ten-fold. REL-2 has been released to sugarbeet researchers because of its superior embryogenic and shoot regeneration abilities for application in biotechnology.

C.J. Tsai and J.W. Saunders. 1999. Encapsulation, germination, and conversion of somatic embryos in sugarbeet. J. Sugar Beet Res. 36:11-32.

ABSTRACT Sugarbeet somatic embryos (SE) of biotech clone REL-2 obtained from callus grown with abscisic acid were experimentally encapsulated with 2% alginate and subsequently germinated and converted into plantlets, in initial efforts necessary for development of artificial seeds. Factors examined were embryo size, alginate companion solution, cold storage duration, and germination substrate. Nonencapsulated SE length category (0.5-1.9, 2.0-2.9, or 3.0-3.9 mm) did not affect germination (GERM) or conversion (CON) rates (87, 89, 87 %, respectively) into complete plantlets on hormone-free Murashige-Skoog (MS) medium. Alginate companion solutions (either hormone-free MS medium or H₂O) had no differential effect on GERM rate (100 %) but did differ in converting embryos to plantlets (81 vs. 64 %, respectively). Subsequent experiments examining cold storage of encapsulated embryos at 4 °C found no lower rate of CONability at 25 °C after 21 days of cold compared with unstored embryos, but after 64 days of

storage at 4 °C, the GERM and CON rates at 25 °C of embryos encapsulated with alginate in MS medium was lower (70 and 45 %, respectively). With alginate in H₂O, respective rates after 64 days of storage at 4 °C were 60 and 20 %. In addition, the GERM rate in soil plates after 64 days cold storage for alginate capsules in MS medium or in H₂O was 38 or 25, respectively. This initial research showed that SE, either nonencapsulated or encapsulated, converted into plants at high frequencies (88 and 81 %, respectively) without cold treatment. Cold storage did not improve the CON rate of encapsulated embryos, but did slow their development. However, these experiments indicated that nonencapsulated and encapsulated embryos were capable of direct GERM after planting on agar plates and in soil.

GERMPLASM REGISTRATIONS

Saunders, J.W., J.M. McGrath, J.M. Halloin, and J.C. Theurer. 1999. Registration of SR94 sugarbeet germplasm with smooth root. Crop Sci. 39:297.

Saunders, J.W., J.C. Theurer and J.H. Halloin. 1999. Registration of EL50 monogerm sugarbeet germplasm with resistance to Cercospora leaf spot and Aphanomyces blackroot. Crop Sci. 39:883.

MASTER OF SCIENCE (M.S.) DISSERTATION (Graduate Advisor: J. W. Saunders):

Goran Srnic, "Inheritance and Intercellular Fluid Protein of a Foliar Disease Lesion Mimic Trait in Sugarbeet (*Beta Vulgaris* L.)" Crop and Soil Sciences, Michigan State Univ., 1999.

Abstract: Disease lesion mimic (DLM) phenotypes in crop plants are characterized by water-soaked spots and lesions on foliage, but close association with forms of disease resistance has been discovered in most such cases. A single DLM sugarbeet from a breeding population was used as a parent in determining the inheritance of the DLM phenotype, using segregation patterns of F_1 , F_2 , F_3 , and BC_1 progenies from a single DLM X wild type cross. Expression of this DLM trait is proposed to be conditioned digenically, by homozygosity of a recessive allele at one locus, and by the simultaneous presence of at least one dominant allele at the second locus (i.e., dlm/dlm_1 : Dlm/-2). DLM occured on older leaves, but was not seen on shoots and plantlets grown on various media in vitro. When intercellular fluid (ICF) proteins were extracted and visualized, defense proteins, including those with chitinase activity, appeared more abundant in leaves from DLM than from wild type plants.

MEETING ABSTRACTS

J.W. Saunders. The Concept of Minimum Assured Frequency of (CMS)-Maintainer Alleles (M.A.F.M.A.) in USDA-ARS Sugarbeet Germplasm Enhancement. 1999 American Society of Agronomy annual meeting, Oct 31-Nov 4, Salt Lake City.

Sugarbeet germplasm from ARS has had direct use potential as parental lines in hybrid cultivars. Most monogerm releases have been Type-O (homozygous for both recessive CMS maintainer alleles x and z), developed only by labor-intensive, calender-consuming identification of Type-O plants, found at 1% or less in most populations, using testcross progeny. Misscoring of testcross progeny occurs in some environments; a 5 degree C difference gave plentiful pollen (25 C) and white anthers (30 C). Recent ARS releases have been less usable as parental lines, as emphasis has shifted to germplasm diversity and combined traits in less finished form. Assurance of CMS maintenance in releases (ie, Type-O) is costly to create, and could be done by the seed industry using ARS releases with pedigree-based minimum assured frequencies of maintainer alleles (ie, M.A.F.M.A.). To that end, population creation and improvement relying on Type-O SP 69550-0 to add higher SUC% to germplasm otherwise high in root smoothness or Rhizoctonia crown and root rot resistance is in progress at East Lansing.

J.W. Saunders. Sugarbeet tissue culture media differentially support the growth of sugarbeet pathogens Rhizoctonia solani, Pythium ultimum, Cercospora beticola, and Aphanomyces cochlioides. 1999 American Society of Sugar Beet Technologists biennial meeting, Feb 10-13, Orlando FL.

Co-culture of pathogen and host plant tissue in vitro offers prospects for studying host defense gene expression, and opportunities for identification and cell selection of resistant genotypes, but pathogen growth characteristics on the medium used to culture the host tissue can determine the feasibility of such systems. Single isolates of sugarbeet pathogens *Rhizoctonia solani* (RZT) and *Pythium ultimum* (PYT) grew well (about 2 cm/day) from mycelial plugs on Murashige-Skoog agar medium with standard 60 mM nitrogen from nitrate and ammonium. *Cercospora beticola* (CER) grew more slowly (about a tenth as fast), and *Aphanomyces cochlioides* (APH) spread rapidly but sparsely. Pathogen growth was also evaluated on nitrogen source variations of MS medium, where the most noteworthy observation was that RZT, PYT and CER grew well with only nitrate as nitrogen source. In general, growth of RZT, PYT, and CER in liquid forms of the media corresponded to growth quantity on the agar versions. APH did not grow at all in liquid MS media with inorganic forms of nitrogen nor with urea, and its sparse growth on corresponding agar media appears due to nitrogenous and sulfurous impurities in the Difco Bacto agar. All pathogens grew to, over, and into sugarbeet tissue cultured on the same plate, leading to host tissue death. CER (due to slow extension growth) and APH (due to sparse growth) should be suitable for future co-culture research with sugarbeet tissue cultures.

GERMPLASM RELEASES

NOTICE OF RELEASE OF **EL52** MONOGERM SUGARBEET GERMPLASM RESISTANT TO RHIZOCTONIA CROWN AND ROOT ROT, APHANOMYCES BLACKROOT, AND CERCOSPORA LEAFSPOT, AND ENRICHED FREQUENCY OF CMS-MAINTAINER ALLELES

The Agricultural Research Service of the U. S. Department of Agriculture, the Michigan Agricultural Experiment Station, and the Beet Sugar Development Foundation announce the joint release of EL52, a sugarbeet germplasm selected for resistance to root-rotting strains (anastomosis group AG-2-2) of *Rhizoctonia solani* Kühn. EL52 was developed at the Sugarbeet and Bean Research Unit, East Lansing, Michigan by the sugarbeet breeding team of Drs. J. W. Saunders, J. H. Halloin. J. M. McGrath, and G. J. Hogaboam (deceased). EL52 also has excellent resistance to Cercospora leaf spot caused by *Cercospora beticola* Sacc. and to blackroot seedling disease and root rot caused by *Aphanomyces cochlioides* Drechs., two of the most destructive sugarbeet diseases in the United States. EL52 is an expected source for development of monogerm parental lines for hybrid cultivars resistant to these three diseases.

EL52 is monogerm, segregates for red and green hypocotyl plants, and is predominantly selfsterile: 6 percent of plants sampled were highly self-fertile. EL52 is enriched for the frequency of the recessive x and z alleles which in the joint homozygous condition maintain male sterility in plants possessing the sterile (S) cytoplasm. EL52 is a bulk of predominantly half-sib seed from 13 of 51 interpollinated plants that had been selected for freedom from disease and for root conformation in the 1997 East Lansing Rhizoctonia crown and root rot nursery. The 51 plants were selected from the six most resistant of 20 half-sib families evaluated in that 1997 nursery. Those 20 half-sib families originated on twenty plants of the half-sib family 85B1-R26 in a crossing block in 1997. 85B1-R26 was one of 25 half-sib families produced in 1985 by interpollinating four tissue culture propagated ramets each of 26 cloned plants. Twenty-five of these plants were from the interrelated East Lansing and Beltsville germplasm pools, and had been selected at East Lansing during 1978-83 from the Rhizoctonia or Cercospora disease nurseries, cloned, and identified as Type-O or near-Type-O individuals. A plant is classified as Type-O or near-Type-O if, by cross to a cytoplasmic-nuclear male sterile tester, its progeny are all, or almost all, respectively, male sterile. Type-O plants have normal (N) cytoplasm and the double homozygous recessive genotype xx zz.

EL52 is moderately resistant to Rhizoctonia crown and root rot, scoring a disease index (DI) equivalent to that of Rhizoctonia resistant specialty cultivar American Crystal Hybrid 1353, but less resistant than resistant checks FC705/1 and FC703 (4.4 compared with 4.4, 3.8 and 3.2, respectively; mean of three readings; DI of 0 = no root rot, and 9 = all plants dead; LSD_{0.05} = 1.1) in the 1998 USDA-ARS evaluation at Ft. Collins, CO. EL52 is resistant to Cercospora leaf spot, receiving a 3.17 disease index compared with 3.25 and 5.33 for the resistant and susceptible checks, respectively (mean of three readings; DI of 0 = no leaf spots and 10 = all plants dead; LSD_{0.05} = 1.23), in the 1998 USDA-ARS evaluation at Ft. Collins, CO. EL52 was not significantly different from SR87 and the two resistant checks (2.4 compared with 2.9, 2.1 and 2.5, respectively; mean of three readings; DI of 1 = full healthy stand and 9 = all plants dead;

 $LSD_{0.05} = 1.17$) in the 1998 Betaseed summer root rot (Aphanomyces) evaluation at Shakopee, MN.

EL52 was tested under the number 98J26-052 where it yielded sucrose concentrations and tons of beets per acre 88 and 107 percent of the mean respectively of the two cultivars ACH555 (American Crystal) and HME17 (Hilleshog-Novartis) in one test at Saginaw, MI in 1998.

EL52 is being released as a germplasm source for breeders to use in developing parental lines combining resistance to Rhizoctonia crown and root rot, Cercospora leaf spot and Aphanomyces seedling disease and root rot. Seed will be maintained by USDA-ARS and is available for use by writing to J. Mitchell McGrath, USDA-ARS, Crop and Soil Sciences Department, Michigan State University, East Lansing, MI 48824-1325. Genetic material of this release has been deposited in the National Plant Germplasm System where it will be available for research purposes, including development and commercialization of new cultivars. It is requested that appropriate recognition be made if this germplasm contributes to the development of a new breeding line or cultivar.

Use of Seed Mixtures of *Rhizoctonia*-Resistant and Susceptible Sugarbeet Varieties for Control of Crown and Root Rot: Effects on Yield and Disease Occurrence.

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Background:

The pattern of disease development typically observed for Rhizoctonia crown and root rot is one where several to many contiguous plants within a row, or within a few adjacent rows are diseased, while plants in other adjacent rows remain nondiseased. This pattern of disease occurrence suggests that *Rhizoctonia* may spread through the soil and surmount the small gap between plants within a row more easily than the larger gap between rows. Occurrences of the disease tend to be widely scattered across fields, making the study of natural disease infestations and their development difficult, in that locations of sites if disease occurrences in a field cannot be predicted in advance of the occurrences.

Rhizoctonia-resistant sugarbeet varieties have become available to Michigan sugarbeet growers in recent years. Although these varieties have yields and sugar concentrations somewhat lower than available Rhizoctonia-susceptible varieties, their use has been advocated for locations where severe crown and root rot problems are anticipated. We proposed that because of the observed pattern of disease development, use of mixtures of seeds from both resistant and susceptible varieties might limit spread of the disease, reducing yield losses from disease, while minimizing yield reductions associated with the resistant varieties. Experiments using seed mixtures of Rhizoctonia-resistant and -susceptible varieties were done in 1998 and 1999 to determine the effects of seed mixtures on yields and disease severity. Because of anecdotal reports that crown and root rot often are severe on fields with no recent history of sugarbeet production, plots were planted at two such locations in 1999.

Methods:

Plantings were done at two locations at which severe crown and root rot was anticipated in 1998 (Hrabal and Terwilligar farms) and at two additional farms (Ivan and Helmrich farms) in 1999. Sites selected as having no recent history of sugarbeet planting were planted in 1999 at the Terwilligar farm and the Bean and Beet Research Farm. Varieties used were the *Rhizoctonia*-

resistant variety RH3, and the susceptible variety E17, and mixtures of the two varieties used contained 1/6, 1/3, and 1/2 RH3. However, at the Helmrich farm (1999), the *Rhizoctonia*-resistant variety C1353 and the susceptible variety C648 were used. Plots were four or six rows, and ran the length of the fields. Each variety or seed mixture was replicated three times in 1998 and four times in 1999 at each location.

Mature beets were harvesed with commercial harvesters, and yields were based upon beets harvested from entire plots. Disease incidences were based on counts of heavily diseased (dead, or with collapsed, necrotic foliage) plants within 600 meters of row within the plots.

Results and Discussion:

Yields (tons per acre and raw white sugar per acre) for plots at the three locations planted with RH3 and E17, and exhibiting crown and root rot are summarized in **Table I**. As anticipated, yields of both beets and sugar were lowest with the *Rhizoctonia*-resistant variety RH3; however, yields were highest with the mixture containing 1/6 (16%) RH3. This yield elevation with the 1/6 mixture of RH3 was statistically different from plots with 100% of either variety alone, when taken across all three locations. Disease occurrences at the three locations planted with varieties RH3 and E17 corresponded closely to percentages of the susceptible variety within plots (**Table II**).

No disease (Rhizoctonia crown and root rot) was observed in 1999 at either of the locations with no recent history of sugarbeet production. Similarly, yields did not show statistically significant differences among treatments at these locations (**Table III**). At the location planted with the varieties C648 (susceptible) and C1353 (resistant), no significant differences were observed among treatments in root yield, raw white sugar per acre, or disease incidence.

We conclude that growers may benefit from reduction of disease severity and enhanced yields when using mixtures of resistant and susceptible varieties containing approximately 1/6 to 1/4 of the resistant variety under conditions where severe crown and root rot is anticipated. Additionally, no statistically important yield penalty was observed when such mixtures were used under apparently disease-free conditions.

Table I. Root yield and raw white sugar per acre of plantings containing either individual sugarbeet varieties resistant (RH3) or susceptible (E17) to Rhizoctonia crown and root rot, or mixtures of the two varieties. Results are the means of observations at two locations in 1998 and one location in 1999 that exhibited losses due to crown and root rot, and are expressed as a percentage of observations for the susceptible variety E17.

Treatment	Ro	ot Yield	RWSA
	_(pe	ercentage	of E17)
100% RH3	-	90.4	82.6
50% RH3+I	E17	99.4	95.3
33% RH3+I	E1 7	100.7	98.6
16% RH3+F	E17	105.3	103.0
100% E17		100.0	100.0

Table II. Root yield and raw white sugar per acre of plantings containing either individual sugarbeet varieties resistant (RH3) or susceptible (E17) to Rhizoctonia crown and root rot, or mixtures of the two varieties. Results are the means of observations at two locations in 1999 that exhibited no losses due to crown and root rot, and are expressed as a percentage of observations for the susceptible variety E17.

			_
Treatment	Ro	ot Yield	RWSA
	(pe	ercentage	of E17)
100% RH3		98.6	93.8
50% RH3+E	17	98.5	98.1
33% RH3+E	17	100.3	98.0
16% RH3+E	17	101.5	101.1
100% E17		100.0	100.0

Table III. Disease occurrence in plantings containing either individual sugarbeet varieties resistant (RH3) or susceptible (E17) to Rhizoctonia crown and root rot, or mixtures of the two varieties. Results are the means of observations at two locations in 1998 and one location in 1999 that exhibited losses due to crown and root rot, and are expressed as a percentage of observations for the susceptible variety E17.

Disease Occurrence
(percentage of E17)
32.1
E17 75.2
E17 70.5
E17 81.6
100.0

Report on a seedling disease survey of Michigan sugarbeet fields, 1999.

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Background:

In recent years there has been increased concern by Michigan sugarbeet growers over declining stand establishment of the crop. While many factors go into the establishment of good stands, such as planting technique, soil structure, weather, seed processing and internal physiology of the seeds themselves, seedling disease is a major determinant. Six main fungal or fungus-like pathogens have been historically linked to seedling mortality in Michigan: Aphanomyces cochlioides, Pythium aphanodermatum, Rhizoctonia solani AG-2-2, R. solani AG-4, Pythium ultimum and Phoma betae. The first four pathogens are most virulent in warm soils and are largely controlled by disease avoidance, planting earlier in the spring so that sugarbeet seedlings grow out of their most vulnerable stage before these pathogens become active. P. ultimum and Phoma betae cause seedling disease at lower temperatures than the others; all commercially-planted sugarbeet seed in Michigan is coated with metalaxyl and thiram fungicides designed to protect against against Pythium spp. and P. betae, respectively. Recently, Pythium strains pathogenic on sugarbeets and resistant to metalaxyl have been isolated from Minnesota sugarbeet fields (Brantner and Windels, 1998). Because of changing cultural practices as well as the potential for pathogen evolution to overcome current control measures, we initiated a disease survey for known sugarbeet pathogens in fields exhibiting stand or seedling disease problems in 1999, and assessed the Pythium isolates for resistance to metalaxyl control methods.

Methods:

<u>Disease survey</u> - Twenty-six fields exhibiting stand establishment or seedling disease problems were identified by Monitor Sugar Co. or Michigan Sugar Co. field personnel. Fields were sampled once in April, May or June, representing the range of planting dates (about 3 weeks to 1 month before sampling) in the 1999 growing season. Seven to ten seedlings with representative disease symptoms were taken from each field. Since field isolations of diseased tissue is sometimes unsuccessful in terms of isolating the causal pathogen, in June, July and August, soil samples were taken from 19 of the 26 fields sampled earlier. Samples consisted of soil immediately below the surface (1-6 cm below surface) dug from several stand gaps, likely to have been sites of seedling disease, which were then pooled into a bulk sample.

Field-sampled seedlings were placed into plastic bags and returned to the lab for further processing. Seedlings were surface-sterilized for 30 seconds in 1% bleach (10% strength of bottle), rinsed 2x in sterile distilled water (dH₂0), and incubated in either 100X20mm culture tubes or on water agar (WA), for purposes of identification. Subsequent transfers were maintained on corn meal

agar (CMA) (Sigma Chemical Co. C-1176) amended with 5mg benomyl/L CMA or 30mg/L CMA metalaxyl to inhibit growth of ascomycetous contaminating fungi or *Pythium* spp. respectively.

Samples of field soil were tested for the presence of seedling pathogens by a bioassay method. Field soil was mixed in a 1:1 ratio with a sterile greenhouse mix consisting of 3:1 sterile field soil:vermiculite to facilitate drainage, and placed into 9 cm diameter round plastic pots. These were planted with 25 sugarbeet seeds (variety: E17) treated with an indicator dye and one of the following fungicide seed treatments: metalaxyl, thiram, metalaxyl + thiram, or no fungicide. Treated seed was kindly supplied by Mr. Kyle Rushing of Gustafson, Inc. The pots were then incubated at either 15 or 25°C (59 or 77°F), placed in individual saucers and watered from below throughout the experiment to prevent crusting. The two temperatures were used to mimic early or later season soil temperatures in the field; the seed treatments were included to prevent "mini-epidemics" of certain pathogens (especially *Pythium* spp.) masking the presence of other pathogens. Two replications of each seed treatment/temperature combination were included. After incubating the pots for 15 days, we harvested any seedlings with disease symptoms. The seedlings were not surface as above, because of concerns that the sterilization procedure eliminated many of the superficially-infecting *Pythium* spp.: seedlings were washed free of soil with tap water, and otherwise treated as outlines above to identify potential pathogens.

Testing for pathogenicity and metalaxyl tolerance of *Pythium* spp. - Since many *Pythium* spp. are saprophytic, including many isolates of species pathogenic on sugarbeets, all cultures of *Pythium* isolated from diseased seedlings were tested for pathogenicity using the methods outlined in Branter and Windels (1998). Briefly, cultures to be tested were grown on 9 cm petri plates of WA (15 g/L; Bacto-Agar, Difco. Co) for 3 days. Then, the entire agar culture was scooped out of the plate was placed inside 9 cm round plastic pots with greenhouse mix (composition as above) and covered with ~5mm of soil. Atop this layer were placed 20 untreated E-17 seeds which were then covered with ~5mm of soil. Pots were incubated and 15 or 25°C, and the cultures were termed pathogenic if seedling stands within each pot differed significantly from blank WA control pots.

Metalaxyl resistance was assessed for all *Pythium* spp., using the methods outlined in Branter and Windels (1998). Isolates with >50% of the growth of non-metalaxyl amended CMA plates on 1ig metalaxyl/L CMA plates were classified as metalaxyl-tolerant.

Results and Discussion:

Disease survey - The most-isolated putative pathogens in April were Pythium spp., but tests of their pathogenicity revealed that only 2 out 12 isolated were pathogenic (Table I). Only 2 out of 10 fields reported pathogenic Pythium spp. One field had two seedlings containing R. solani. A range of other fungi such as Papularia, Stemphyllium and Alternaria were isolated (data not shown) usually classified as saprophytes or weak pathogens. It seemed likely that most of the fungi isolated from seedlings in April were "symptoms" rather than causes of disease; seedlings were weakened or damaged by other factors such as the mild frost which hit the sugarbeet growing area in mid -April of 1999. However the isolation method (which included a surface-sterilization) may have excluded certain Pythium spp. and certainly the small sample size in any particular field may have underestimated (or overestimated) the relative importance of a particular pathogen. In testing of soil

samples from the same fields (Table II, fields 429-2 through 429-10) *Pythium* spp. were isolated from diseased seedlings in soil bioassays incubated at 15°, and a range of pathogens (*Pythium* spp, *A.cochlioides* and *R. solani*) were isolated at 25°C. Since soil sampled were taken from stand gaps, it is possible that the presence of these warm-soil pathogens may have exacerbated the stand problems in these fields.

In May and June, many more pathogenic *Pythium* spp. were isolated from the fiels, as well as some A. cochlioides and R. solani (**Table I**) which would be expected as warm soil temperatures are more conducive to seedling diseases caused by these pathogens.

In the soil bioassays, incubation at 15°C favored the isolation of *Pythium* spp. from diseased seedlings over *A. cochlioides* or *R. solani*; interestingly, *Aphanomyces cochlioides* was isolated from seedlings incubated at 15°C in fields 429-4 and 623-43 (**Table II**). Work is underway to assess the temperature optima in terms of pathogenicity of these isolates. In all cases, fewer damping-off symptoms were seen at 15°C than 25°C; however, emergence counts were generally lower (data not shown). It may be possible that more pre-emergence damping-off occurred in 15°C soils. The pathogens isolated from these two incubation temperatures roughly mirrored the pathogens isolated from the field in the cooler soils of April and the warmed soils of May and June.

Metalaxyl resistance of Pythium spp. isolates – The presence of metalaxyl tolerance was limited to three (and possibly four) of the fields sampled. However it was strongly correlated to pathogenicity: of the 12 pathogenic *Pythium* isolates recovered, 10 were metalaxyl tolerant. No pathogenic isolates of *Pythium* were recovered from one field, 609-28 (a field near an irrigation pond at the Bean and Beet Research Farm, St Charles, MI, a field with a history of heavy seedling mortality) but soil testing recovered many *Pythium* isolates from pots with seed treatments containing metalaxyl (Table II). More work is necessary to assess their resistance to metalaxyl in vitro.

In the soil bioassay experiment, *Pythium* spp. were isolated, in general, in the untreated or thiram-only experiments, except in fields429-3, 609-28, 623-43 and 623-44 (**Table II**).

Future sampling will continue in 2000, with additional soil testing of problem fields with the goal of developing a "Seeding Disease Potential" assay designed to forecast potential seedling disease problems with either a early or late-planting regime. Field sampling of seedlings will also be continued. Because of the low success rate of isolating known sugarbeet pathogens from symptomatic tissue, additional samples will be taken, and the pathogenicity of putative secondary colonizers of diseased tissue such as *Fusarium* and *Alternaria* to sugarbeet seedlings will be assessed, to gauge if virulence to sugarbeet seedlings has developed in these fungi.

Reference:

Brantner JR and CE Windels. Variability in sensitivity to metalaxyl in vitro, pathogenicity, and control of *Pythium* spp. on sugar beet. Plant Disease. 82(8) pp. 896-899.

Table I: Number of known sugarbeet pathogens isolated from seedlings with disease symptoms sampled in April, May and June (early through late plantings) of 1999. Number of metalaxyl-tolerant Pythium isolates in parentheses.

Pyth	ium spp.	Aphanomyces cochlioides	Rhizoctonia solani
Pathogenic (Metalaxyl tolerant)	Nonpathogenic (Metalaxyl tolerant)		
April			
2 (2)	10(0)	0	2
May			
7(6) 10(2)		3	2
June			
2(0)	3	4	5

Table II: Known sugarbeet pathogens isolated in soil bioassay experiment; sorted by code of field isolated from; temperature incubated (15°C or 25°C) and seed treatment (N=None; M=metalaxyl; T=thiram; M+T=metalaxyl+thiram). Pyth = Pythium spp., Rhiz=Rhizoctonia solani, Aph=Aphanomyces cochlioides

	15∘C				25∘C			
Field	N	M	T	M+T	N	M	T	M+T
429-2	Pyth				Aph	Aph Rhiz	Aph Rhiz	
429-3					Pyth	Aph Pyth		Aph Pyth
429-4	Pyth	Aph			Pyth		Pyth	Aph
429-5						Rhiz		
429-6	<u> </u>							
429-9								
429-10					Pyth Rhiz	Aph Rhiz	Aph Rhiz	Aph Rhiz
521-15	Pyth				Pyth Rhiz		Pyth	
521-16			1		Pyth			
521-17								
521-18					Pyth	Rhiz		
521-19			Pyth				Pyth	
604-31								
604-32					Pyth			
604-33					Pyth		Aph Pyth	
608-25								
609-28	Pyth	Pyth		Pyth	Pyth	Pyth	Pyth	Aph Pyth
623-42					Pyth		Pyth	Aph Pyth Rhiz
623-43	Pyth	Aph Pyth	Aph		Rhiz	Rhiz	Pyth	Aph Pyth

SUGAR BEET RESEARCH

1999 REPORT

Section F

Texas Agricultural Experiment Station Bushland, Texas

Dr. C. M. Rush, Professor

Cooperation:

Holly Sugar Corporation - Sugar Land, Texas Western Sugar Company - Denver, Colorado

This research was supported in part by funds provided through the Beet Sugar Development Foundation (Projects 503, 506 and 507)

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by C. Rush		F3

Interactions Between BNYVV and BSBMV

Charlie Rush Bushland, Texas

In previous studies we have shown that BSBMV and BNYVV are wide spread in most sugar beet growing regions of the United States. The two viruses are often found together in the same fields and sometimes infecting the same beet. Most ELISA methods of virus detection are comparable to results obtained using Western Blots but in cases where the most accurate test results possible are required, molecular techniques, such as PCR, should be used. However, molecular tests are not appropriate for routine diagnostics because of expense and time requirements. Because of the similarities between BNYVV and BSBMV, cross-reactions in some ELISA tests are possible if test conditions are not suitably stringent.

BNYVV and BSBMV are very closely related at the molecular level and have the same genomic organization. The same soilborne fungus vectors the two viruses and conditions for disease development are similar. We have shown that BSBMV can cause significant damage to sugar beets, especially under conditions of high soil moisture. However, the greatest concern is the possibility of recombination between BNYVV and BSBMV. Recombination is relatively common between RNA viruses and is a recognized method that viruses use to develop new strains that can differ from the two "parents" in virulence. Although good disease resistance to BNYVV is available, it was unknown for sure whether BNYVV resistance genes also conferred resistance to BSBMV.

This year, we conducted studies to evaluate interactions between BNYVV and BSBMV at the field level. We conducted studies to verify the susceptibility of BNYVV resistant germplasm to BSBMV and also screened the core collection of *Beta maritima* for accessions with resistance to BSBMV. In addition, we conducted field studies to evaluate the effect of various irrigation rates on soilborne diseases of sugar beet caused by various soil fungi and soilborne viruses.

Methods

Interactions between BNYVV and BSBMV- A survey was made in Colorado and Minnesota for fields with BNYVV, BSBMV, or both viruses. Based on the results of an initial survey several fields were grid soil sampled. Selected fields were marked off in 60, one-acre grids and soil samples taken from each grid cell. Samples were geo-referenced for future identification. In addition to these grid soil samples, intensive sampling was conducted in several fields by taking grid samples on a 10' grid pattern. Soil samples were taken to the laboratory and bioassays were initiated by planting seed in individual samples. All soil samples from an individual field were planted at the same time, and field samples were planted approximately every six weeks. Plantings were staggered to allow time for sample processing after harvest, which is approximately 10 weeks after planting. After harvest, plants were tested by ELISA to determine the distribution of BSBMV and BNYVV in the field and by SSCP analysis to determine the degree of genomic variability of BSBMV in fields and the possibility of recombination. In one field, plant samples, in addition to soil samples, were taken from one of the intensively sampled fields. The intensive sampling in this field was in a BNYVV disease-screening nursery.

BNYVV germplasm resistance to BSBMV - In 1998, a rhizomania resistance cultivar nursery was sampled and we found that the BNYVV resistant cultivars seemed to be highly susceptible to BSBMV. In 1999, a study was initiated to determine whether cultivars with genetic resistance to BNYVV were susceptible to BSBMV. Twenty entries with varying levels of resistance to BNYVV, ranging from 0-100%, were grown in a field naturally infested with BSBMV and BNYVV. Two times during the season, samples were collected and tested by the ELISA test. Absorbance values were compared between the susceptible and resistance lines and correlation analysis was conducted to determine whether there was any association between absorbance intensity and degree of resistance.

Screening the Beta maritima core collection for resistance to BSBMV - The USDA Beta germplasm collection is maintained in Pullman, Washington. The core collection of Beta maritima, which contains approximately sixty accessions from around the world, was obtained from the collection curator and screened for resistance to BSBMV. Seed from each accession were planted in soil infested with BSBMV and then the plants were grown, under conditions conducive for virus infection, for ten weeks. After this period of baiting, plants were harvested and tested by ELISA for infection by BSBMV. There were seven replications of each accession in each of two separate tests.

Irrigation Study: Sugar beet varieties Kojak and Ranger were planted at a rate of 7 seeds per foot on April 1, 1999. Irrigation was supplied by a center pivot irrigation system, with 60" drops equipped with LEPA nozzles. Three irrigation treatments were implemented during the growing season (2.5" every week, 2.5" every other week and 5.0" every third week). On September 9, plots were harvested. Sugar beets were topped, weighted, given a disease rating, and percent sucrose was determined.

Results

Interactions between BNYVV and BSBMV- To date, three sets of soil samples have been planted and one harvested. The intensively sampled field in which plant samples were taken was the first field to be tested. Plant samples collected from the variety test displayed obvious systemic symptoms of BSBMV and typical symptoms of BNYVV infection also, but when the collected root samples were tested by ELISA, no samples tested positive for either BSBMV or BNYVV. The test was repeated and again no positive samples were obtained. Because of the prevalence of systemic symptoms on the plants that were sampled, we tested the plants a third time, but used the Western Blot test. Results were positive and 96% of the plants tested positive for BSBMV but less than one percent tested positive for BNYVV alone. Sixteen percent tested positive for both BNYVV and BSBMV. When we harvested bioassay plants from the soil samples taken from the same plots, results were opposite those of the field grown plants tested by Western Blot analysis. Thirty eight percent of the bioassay plants tested positive for BNYVV and but only two percent tested positive for BSBMV.

Results of these studies seem contradictory, but in fact they are not and they provide an important hint to the interactions between BSBMV and BNYVV. It appears from these results that BNYVV infects first during the season but by the end of the season, BSBMV has dominated in the competition. This explains why BSBMV was the predominant pathogen in the field sampled beets, harvested at the end of the season, but BNYVV was predominant in the 10-

week-old bait plants. The results of this study support previous greenhouse studies where BSBMV was dominant over BNYVV in dual infection studies.

BNYVV germplasm resistance to BSBMV - Results of this year's repeated study, corroborated those from a preliminary study conducted in 1998. Cultivars with resistance to BNYVV are totally susceptible to BSBMV and there is no correlation between the degree of resistance to BNYVV and the virus titre of BSBMV in infected plants (Table 1). Since most genetic resistance to BNYVV is, at present, based on the Holly resistance gene, it is unlikely that any BNYVV resistant cultivars will possess significant resistance to BSBMV. Therefore, if recombination between BSBMV and BNYVV occurred, current varieties resistant to BNYVV might become susceptible. Furthermore, all current sugar beet varieties are susceptible to BSBMV and a particularly virulent isolate could cause significant disease loss. Even a mild isolate, in the presence of extremely wet soil conditions, might cause significant losses in quality and root yield.

Table 1. 1999 BNYVV/BSBMV Susceptibility Study

Cultivar	% BNYVV Resistance	O.D.	ANOVA
Beta 4035 R	90	0.354	NS
Beta 4038 R	90	0.229	NS
Beta 4006 R	90	0.579	NS
Beta 1399	0	0.597	NS
Maribo 9372	0	0.08	NS
Seedex 705	75	0.271	NS
Kojak	50	0.096	NS
Monohy 970601101	100	0.085	NS
Monohy 9706035201	100	0.316	NS
Monohy 9706047501	50	0.623	NS
Monohy 9706047601	75	0.089	NS
Monohy 9706047801	75	1.009	NS
Monohy 9706048201	50	0.308	NS
Monohy 9706048301	50	0.105	NS
Monohy 9804011001	0	0.51	NS
Monohy 9806003701	100	1.15	NS
Monohy 9155	0	1.351	NS
Monohy 9255	0	1.183	NS
Monohy RH3	0	0.093	NS
Monohy 1639	100	0.279	NS

Correlation Coefficient = -0.14 NS

Screening the Beta maritima core collection for resistance to BSBMV - Approximately 60 accessions from the Beta maritima core collection were screened for resistance to BSBMV. Two accessions were identified, 546417 from France and 546404 from the Netherlands, that appear to have significant resistance to the virus. In both accessions, none of the test plants tested positive for BSBMV in the two repeated tests. There were seven replications in each test so the odds of negative results due to escape from infection are minimal.

Irrigation Study - Environmental conditions were particularly wet this year, making this a difficult year for imposing differential irrigation treatments. Only in the later part of the growing season we were able to impose a limited irrigation treatment that had a significant effect in controlling disease incidence. Table 2 summarizes the results obtained from the study. Regardless of the irrigation treatment Kojak had significantly higher disease incidence than Ranger. Both varieties, however, had less disease when grown under limited irrigation. Although no significant yield differences were found between the two varieties, Ranger had the tendency to yield more than Kojak. Percent sugar, on the other hand, was significantly higher in Ranger under full irrigation. Limited irrigation did not reduce yield or percent sugar in either variety.

Table 2.

Variety	Full Irrig	ation		Limited Irrigation			
	Disease Rating	Yield Tons/ac	Sugar %	Disease Rating	Yield Tons/ac	Sugar %	
Kojak	4.0 A a	19.4 A a	13.7 A a	2.4 A b	17.4 A a	14.3 A a	
Ranger	2.6 B a	26.8 A a	14.2 B a	1.4 B b	23.6 A a	14.4 A a	

Means followed by the same upper case letter within a column are not significantly different. Means followed by the same lower case letter within a row are not significantly different.

Discussion

Results of the field grid sampling study verified that interactions occur between BNYVV and BSBMV and that BSBMV becomes dominant over BNYVV as the season progresses. However, our greenhouse study suggests that BNYVV infects plants first (it may be able to infect plants at a cooler soil temperature) and therefore is able to cause severe disease even though BSBMV eventually dominates the infection. We have not completed SSCP analysis of the samples yet and do not know whether the interaction between BNYVV and BSBMV includes recombination or not

If recombination does occur between BNYVV and BSBMV it is impossible to predict whether a hybrid strain that can infect BNYVV resistant cultivars and cause significant disease will develop or not. To date, all BNYVV resistant cultivars tested have been susceptible to BSBMV. Nearly all these cultivars possess the Holly resistance gene, so our base for genetic resistance is very narrow. We know that recombination is common among RNA viruses and since BNYVV and BSBMV are so genetically similar and are often together in a single infection in the field, recombination is likely instead of just possible. As a precaution, breeders should begin to identify and incorporate BSBMV resistance into breeding lines for quick introduction into cultivars. Results of our BSBMV screening study are encouraging because two and possibly three *Beta maritima* accessions tested negative for BSBMV infection in two well replicated studies. It will be important to determine whether these accessions have any affect on infection by BNYVV.

The field where the irrigation study was conducted was infested by numerous soilborne pathogens, including Rhizoctonia, Fusarium, Aphanomyces, BNYVV and BSBMV. Reduced irrigation reduced disease incidence that resulted in yields equal to those in higher irrigation plots. However, Ranger, a BNYVV susceptible cultivar, yielded better than Kojak, which is resistant to BNYVV. This demonstrates that when a BNYVV tolerant cultivar is grown in a field infested with multiple pathogens, tolerance to BNYVV may be of secondary importance to the other pathogens in the field. BNYVV resistant cultivars are the best means of managing rhizomania but growers must be prepared to use additional disease management strategies if the field is infested with more than BNYVV.

SUGARBEET RESEARCH

1999 Report

Section G

Molecular Plant Pathology Laboratory

Agricultural Research Service

United States Department of Agriculture

Beltsville, Maryland

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- Mujer, C.V. and A.C. Smigocki. Wound-inducible cytochrome P-450 from *Nicotiana plumbaginifolia*. Physiol. Plant. (submitted)
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- Smigocki, A.C., S. Heu, C. Wozniak and G. Buta. Leaf extracts from transgenic tobacco that express the cytokinin biosynthesis gene ipt are lethal to <u>Manduca sexta</u> L. (Lepidoptera: Sphingidae) and <u>Tetanops myopaeformis</u> von Roder (Diptera: Otitidae). J. of Economic Entomology (submitted 10/99).
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BIOTECHNOLOGICAL STRATEGIES FOR EFFECTIVE CONTROL OF THE SUGARBEET ROOT MAGGOT (TETANOPS MYOPAEFORMIS RODER).

Smigocki^{*}, Ann¹, Stephen Wilhite², Tom Elden², Scott Armstrong³ and Chris Wozniak^{1,4}, Molecular Plant Pathology Laboratory, ²Soybean and Alfalfa Research Laboratory, ARS, USDA, Beltsville, MD 20705, ³Department of Entomology, North Dakota State University, Fargo, ND 58105 and ⁴Biopesticide and Pollution Prevention Division, US Environmental Protection Agency, Washington, D.C. 20460

Two approaches are being undertaken for management of the most devastating pest of sugarbeet in the US, the sugarbeet root maggot (SBRM). One approach involves the expression in transgenic sugarbeet plants of proteinase inhibitor genes which have specific activity against the root maggot's digestive proteases. These enzymes are essential for the release of nutrients for normal growth and development. Extracts of midguts excised from feeding second instar larvae were analyzed for specific protease classes using an inhibition assay. More than 86% of the gut protease activity was inhibited by 2 mM phenyl methyl sulfonyl fluoride, a serine protease inhibitor. Less than 3% inhibition was observed with 50 μM E-64, a cysteine protease inhibitor, and no inhibition with Pepstatin A, an aspartyl protease inhibitor. Using azocasein as a substrate, maximum protease activity was detected at pH 8.5, consistent with the serine class of proteases. Another approach being evaluated is the effect of cytokinin-induced insecticidal compounds on the SBRM larvae. A 1% suspension of leaf surface extracts from Nicotiana plumbaginifolia plants transformed with a cytokinin biosynthesis gene induced a twitching response and death of 30% of the first instar SBRM larvae at 72 hr. After 120 hr, 92% of the larvae were dead as compared to about 25% of the controls. Sugarbeet plants transformed with the cytokinin biosynthesis gene fused to a woundinducible or a tuber-specific promoter have been regenerated for further analysis of the effect of cytokinins on defense responses.

CARBOHYDRATE CONTENT OF SUGARBEET (BETA VULGARIS L.)
TRANSFORMED WITH A CYTOKININ BIOSYNTHESIS GENE. Snezana Ivic¹, Iris McCanna¹, Richard Sicher² and Ann Smigocki¹ Molecular Plant Pathology Laboratory, ²Climate Stress Laboratory, ARS, USDA, Beltsville, MD.

To study the role of cytokinins in carbon partitioning, sugarbeet clone Rel-1 was transformed with the isopentenyl transferase *ipt* gene fused to a wound-inducible proteinase inhibitor II (Pin2) or a tuber-specific patatin (Pa) gene promotor. Two transformation methods were used, *Agrobacterium*-mediated cotyledon transformation and particle bombardment of embryogenic hypocotyl callus. For root initiation, transformed shoots had to be exposed to high auxin concentrations (50 mg IBA/I) for 24 hours as compared to normal shoots that were maintained on 3 mg IBA/I. *Ipt* shoots rooted in 4-8 weeks and the controls in 2 weeks. All *ipt*-transformed plants exhibited phenotypic characteristics associated with elevated cytokinin levels. Some showed

increased adventitious shoot formation while others had reduced apical dominance, a large, proliferative crown and a very small root mass. Others exhibited slower growth and an overall reduction in the number and size of leaves. Leaf and taproot cytokinin levels were up to 17 and 2 times higher, respectively, than in normal plants. In one transformant, about a 9 fold increase in leaf sucrose levels was observed while the glucose content was 18 times higher. No corresponding increase in sucrose and glucose levels was observed in the taproots of this plant.

INHIBITION OF ASPARTYL AND SERINE PROTEINASES IN THE MIDGUT OF SUGARBEET ROOT MAGGOT WITH BIOCHEMICAL AND PLANT-DERIVED PROTEINASE INHIBITORS. Stephen E. Wilhite¹, Thomas C. Elden¹, Borut Strukelj², Scott Armstrong³, and Ann C. Smigocki⁴ ¹Soybean and Alfalfa Research Laboratory, Agricultural Research Service, U.S. Department of Agriculture, Beltsville, MD 20705, USA, ²Department of Biochemistry and Molecular Biology, Jozef Stefan Institute, Jamova 39, SI-1000, Ljubljana, Slovenia, ³Plant and Soil Sciences Department, Texas Tech University, Lubbock, TX 79409, USA, ⁴Molecular Plant Pathology Laboratory, Agricultural Research Service, U.S. Department of Agriculture, Beltsville, MD 20705, USA.

The use of genes encoding proteinase inhibitors (PIs) to transform crop plants for resistance to insect pests (see Jouanin et al., 1998, and; Schuler, et al., 1998, for reviews) may represent an alternative approach to insect control. PIs occur naturally in a number of plant species and are likely a part of the natural defense mechanism against insects (Green & Ryan, 1972). PIs specifically bind and inhibit the action of digestive proteinases in the insect midgut, thereby exerting a deleterious effect on insect growth and development (Jongsma & Bolter, 1997, for review). Due to significant variation in the types and properties of proteinases utilized by insects for dietary purposes (see Terra & Ferreira, 1994, for a review), and the altered specificity that plant PIs possess toward such proteinases (Keilova & Tomasek, 1976; Abe et al., 1994; Brzin et al., 1998; Christeller et al., 1998; Pernas et al., 1998), it is necessary to characterize the proteolytic activities of each individual pest species in order to devise a rational control strategy. The present study examines the effect of pH, low-molecular weight inhibitors, and plant-derived PIs on general substrate hydrolysis to identify the major midgut proteinases of the SBRM.

WOUND-INDUCIBLE CYTOCHROME P450 FROM <u>NICOTIANA</u>
PLUMBAGINIFOLIA. Cesar V. Mujer and Ann C. Smigocki Molecular Plant
Pathology Laboratory, Agricultural Research Service, U.S. Department of Agriculture,
Beltsville, MD 20705, USA.

Two Nicotiana plumbaginifolia cDNA clones, CYP72A2 and npl2, with high sequence similarity to cytochrome P450 monooxygenases were isolated using reverse transcription-polymerase chain reaction. CYP72A2 has an open reading frame of 1524 nucleotides and its deduced 508 amino acid sequence has 45% identity to Catharanthus

roseus P450 CYP72A1. npl2 is similar to CYP72A2 except for an 82-nucleotide deletion within its coding region and an internal stop codon. Southern blot analysis indicated that there are at least three copies of the CYP72A2 gene and that they are induced by mechanical wounding, insect chewing (Manduca sexta) and cytokinin application. In N. plumbaginifolia plants transformed with a wound-inducible cytokinin biosynthesis gene construct (PI-II-ipt), mechanical wounding of the leaves induced a 6-fold increase of CYP72A2 messages at 6 h in comparison to a 2-fold induction after 12 h in wounded, untransformed leaves. A similar response was observed when plants were sprayed with 5 x 10⁻⁵ or 5 x 10⁻⁶ M zeatin or when M. sexta larvae fed on the leaves. The response to feeding larvae and wounding was systemic. Using polyclonal antibodies raised against three internal regions of the deduced CYP72A2 protein, a 58.8 kDa polypeptide was detected in leaves of N. plumbaginifolia as well as in the leaves of 4 other plant species. The modulation of CYP72A2 expression by cytokinins and the possible role of P450 in plant defense responses are discussed.

INHIBITION OF CYSTEINE AND ASPARTYL PROTEINASES IN THE ALFALFA WEEVIL MIDGUT WITH BIOCHEMICAL AND PLANT-DERIVED PROTEINASE INHIBITORS. Stephen E. Wilhite¹, Thomas C. Elden¹, Joze Brzin², and Ann C. Smigocki³ Soybean and Alfalfa Research Laboratory, Agricultural Research Service, U.S. Department of Agriculture, Beltsville, MD 20705, USA, ²Department of Biochemistry and Molecular Biology, Jozef Stefan Institute, Jamova 39, SI-1000, Ljubljana, Slovenia, ³Molecular Plant Pathology Laboratory, Agricultural Research Service, U.S. Department of Agriculture, Beltsville, MD 20705, USA.

Proteolytic activities in alfalfa weevil (Hypera postica) larval midguts have been characterized. Effects of pH, thiol activators, low-molecular weight inhibitors, and PIs on general substrate hydrolysis by midgut extracts were determined. Hemoglobinolytic activity was highest in the acidic to mildly acidic pH range, but was maximal at pH 3.5. Addition of thiol-activators DTT, 2-ME, or L-cysteine had little effect on hemoglobin hydrolysis at pH 3.5, but enhanced azocaseinolytic activity two to three-fold at pH 5.0. The broad cysteine proteinase inhibitor E-64 reduced azocaseinolytic activity by 64% or 42% at pH 5 in the presence or absence of 5 mM L-cysteine, respectively. Inhibition by diazomethyl ketones, Z-Phe-Phe-CHN₂ and Z-Phe-Ala-CHN₂, suggest that cathepsins L and B are present and comprise approximately 70% and 30% of the cysteine proteolytic activity, respectively. An aspartyl proteinase component was identified using pepstatin A, which inhibited 32% (pH 3.5, hemoglobin) and 50% (pH 5, azocasein) of total proteolytic activity. This activity was completely inhibited by an aspartyl proteinase inhibitor from potato (API), and is consistent with the action of a cathepsin D-like enzyme. Hence, genes encoding PIs with specificity toward cathepsins L, B and D could potentially be effective for control of alfalfa weevil using transgenic plants.

Gene Transfer to Optimize the Sucrose Storage Capacity of the Sugarbeet Taproot

BSDF Project 810

Ann C. Smigocki

Changes in phytohormone profiles of sugarbeet taproots between sowing and harvest have been determined and related to initiation of cambia, cell division of the cambia and rapid cell expansion stages in root development. It is well established that cytokinins induce cell division and, in taproot-derived sugarbeet suspension cultures, cytokinin levels were shown to peak just prior to cytokinesis. These results suggest that higher cytokinin levels in the taproot will lead to increased cell division, additional vascular rings and increased sucrose yield. In addition, cytokinins have been identified as having functional significance in the control of assimilate movement in plants, particularly by altering phloem unloading, sink initiation, and sink strength and capacity.

Higher endogenous cytokinin levels are anticipated to increase the sink activity of the taproot leading to an increase in the overall root productivity and a decrease in the leaf sucrose storage pools. The removal of more sucrose at the source may also decrease the feedback inhibition on the system and might be expected to increase the maximum rate of photosynthesis. Additionally, increasing cell division and the number of vascular rings in the taproot is expected to produce a low-tare sugarbeet with globe-shaped storage root with fewer branches or grooves. A low-tare sugarbeet would be of great benefit to the farmers, processing plants and the environment.

Since field applications of phytohormones are of limited value due to high costs and rapid degradation, we have genetically engineered sugarbeets for production of high cytokinin

levels in the taproot. To increase endogenous cytokinins in the taproot, a bacterial cytokinin biosynthesis gene ipt was fused to a tuber-specific promoter from the patatin gene of potato and introduced into sugarbeet using the method of particle bombardment of embryogenic hypocotyl and cotyledon callus. Leaf zeatin riboside concentration in two independent transformed lines was up to 18-fold higher than in the control, while a corresponding 2-fold increase was observed in the taproots. Elevated cytokinin levels were associated with distinguishable morphological alterations that are commonly seen in ipt transformants, i.e. reduced root growth and leaf surface area and adventitious shoots development. Leaf concentrations of major carbohydrates, sucrose, glucose and starch were not significantly different from the control plants. Taproots of mature (8-12 month) transgenic plants were greatly reduced in size and had lower carbohydrate concentrations as compared to the controls. However, sucrose concentrations in young (5 month) taproots of two of the transformants were elevated in comparison to the untransformed control. These preliminary results support the hypothesis that higher cytokinin levels may enhance sucrose accumulation in younger taproots but become detrimental to normal development as the plant matures.

Engineering sugarbeets with multiple proteinase inhibitor genes for enhanced tolerance to the sugarbeet root maggot

BSDF Project 811

Ann C. Smigocki

The sugarbeet root maggot (SBRM) *Tetanops myopaeformis* von Röder (Diptera: Otitidae) was first described as a sugarbeet pest in Utah in the 1920's. It is now considered the major sugarbeet pest of the central and western sugar-beet-growing areas in the United States and Canada. More than half of the U.S. sugarbeet fields are infested. Developing SBRM larvae feed on roots throughout the growing season, inflicting significant crop damage and yield losses as high as 23%. Control has come primarily through the application of pesticides to sugarbeet fields in order to reduce larval populations. Cultural control practices, such as crop rotation, are made difficult by the mobility of the adult flies, and the existence of several weed species as alternate hosts hinders population control. Currently no biological control measures are available. In the next few years all chemical pesticides effective against the maggot will likely be removed from EPA approved registrations. Therefore, an urgent need exists to develop effective, environmentally safe approaches to target this pest.

The use of genes encoding proteinase inhibitors to transform crop plants for resistance to insect pests represents an alternative approach to insect control. Proteinase inhibitors occur naturally in a number of plant species and are likely a part of the natural defense mechanism against insects. Proteinase inhibitors specifically bind and inhibit the

action of digestive proteinases in the insect midgut, thereby exerting a deleterious effect on insect growth and development. Due to significant variation in the types and properties of proteinases utilized by insects for dietary purposes, and the altered specificity that plant proteinase inhibitors possess toward such proteinases, it is necessary to characterize the proteolytic activities of each individual pest species in order to devise a rational control strategy.

Latest studies on inhibition of insect protease activities by proteinase inhibitors indicate that a combination of inhibitors incorporated into insect diets is more toxic at levels where individual inhibitors are not toxic. In addition, higher levels of more than one proteinase inhibitor have been found in insect resistant vs. susceptible plants. Therefore, introduction of multiple proteinase inhibitor genes into transgenic plants will most likely prove to be the most effective and perhaps sustained means of controlling insect infestations.

We examined the effect of pH, low-molecular weight inhibitors, and plant-derived proteinase inhibitors on general substrate hydrolysis to identify the major midgut proteinases of the SBRM. We dissected out midguts from feeding second instar sugarbeet root maggot larvae that were collected in St. Thomas, ND in the summer of 1998. Major classes of digestive proteinases were identified. Proteolytic activity in larval gut extracts peaked at pH 2.5 and 9.5. Addition of low-molecular weight biochemical inhibitors targeting three major classes of insect digestive proteinases revealed that Pepstatin A, an aspartly proteinase inhibitor, was by far the most effective inhibitor at pH 3.0 (83.9% inhibition). A cysteine proteinase inhibitor, E-64, which has high potency toward virtually all known cysteine proteinases had only minor inhibitory activity (6.5%). At pH

8.5, treatment with PMSF inhibitor resulted in a sizable decrease in proteolysis (47.3% inhibition) suggesting that serine proteinases are major contributors to proteolysis at the higher pH. Proteinase inhibitors purified from plants were also tested. Squash aspartyl proteinase inhibitor (SQAPI) blocked virtually all the proteolytic activity at pH 3.0, thus confirming the importance of the aspartyl class at acidic pH. Soybean trypsin-chymotrypsin inhibitor (Bowman-Birk inhibitor I) blocked nearly all proteolysis at pH 8.5, suggesting the presence of trypsin and/or chymotrypsin-like serine proteinases in the extract. Overall, our results indicate that majority of the digestive enzymes found in the actively feeding maggot midguts are aspartyl and serine proteases with a relatively small portion of the activity being associated with cysteine proteases.

Bioengineering Sugar Beets for Disease Resistance

L.David Kuykendall

Molecular Plant Pathology Lab, Beltsville, MD

Sugar beets, long regarded as recalcitrant for both DNA transformation and plant regeneration from individual transformed cells, two essential prerequisites for biotechnological improvement of the crop, could theoretically benefit from 21st century science. Snyder, Ingersol, Smigocki & Owens (1999) reported transgenic sugarbeets carrying *ipt*, an agrobacterial cytokinin gene, which may influence insect pest susceptibility and ones encoding antimicrobial peptides that could enhance resistance to pathogens.

Using methodology as described in the 1998 BSDF Annual Report, we have since identified two transgenic sugar beet clones as candidates for having improved leafspot resistance due the expression of introduced antimicrobial protein gene(s). Unlike all of the other clones examined, they have some ability to inhibit the growth of *Cercospora beticola*, the fungus that causes leafspot disease. In the majority of the U.S. sugar beet -growing acreage, leafspot reduces both yield and sucrose percentage by as much as one third or more. Reverse transcriptase polymerase chain reaction, called RT/PCR, is being used to measure the expression of the introduced genes. For determining either the biological or potential agronomic significance, these new sugar beet genotypes have been vegetatively propagated with the plan of examining a number of healthy greenhouse-grown plants for their ability to resist foliar germination of *C. beticola* spores.

Supported in part by this BSDF project, Joe Saunders of MSU, East Lansing, MI, has recently visited Beltsville to transfer methodologies for clonal selection and regeneration.

Growing out of this was a newly devised method of incubating embryogenic callus of sugar beet

under light on a rotary shaker in medium containing the appropriate growth regulator and μM quantities of cercosporin to select for resistance. Toxin resistant shoots can develop in 14 days.

Since only a low efficiency of transformation and regeneration is available and since resources are limited, we must therefore carefully choose which bioengineered, disease-resistance-conferring genes to introduce into sugar beet. Dr. Bob Davis, RL of MPPL, has obtained Agriculture Research Service funding to support a postdoctoral researcher to use viral vectors to study gene expression in transfected sugar beet lines, a new approach deserving careful attention.

Since 1998, when this project was first initiated as BSDF #830, a new approach actively underway at Beltsville, Maryland has been aimed at the genetic transfer into sugar beet of a bioengineered cfp gene from Cercospora generously supplied by Greg Upchurch of Raleigh, NC. The laboratory construct we have generated places cfp under the control of the stress-inducible Ubi7 promoter from potato, courtesy of Bill Belknap of Albany, CA. Following the suggestion of Dr. Roger Lawson, we needed an ARS-owned non-proprietary promoter and Ann Smigocki suggested Ubi7. Belknap provided the 1.7kb Ubi7 promoter (NCBI accession number stu26813) as a gus reporter gene fusion in pUC19, an E.coli plasmid vector. BamH1 restriction enzyme digestion was used to liberate the intron-carrying (prevents expression in bacteria) Ubi7 promoter from both the gus fusion and from the pUC19 vector. Purified restriction fragment was ligated to cfp (NCBI accession number AFO91042) carried on pBS, another E. coli cloning vector, and then the ligation mix was used to transform competent cells of E.coli strain DH5a. Among the transformants, clones carrying the Ubi7 promoter fused immediately upstream of the cfp gene were identified and then the exact orientation of the insert DNA was experimentally ascertained by multiple enzyme restriction analysis. Now, the desired gene fusion are being inserted into an Agrobacterium Ti-based vector so that transformation of sugar beets can then be performed. Ann

Smigocki plans to cooperate in the latter stages. Since the cell-killing, lesion-forming cercosporin toxin is presumed with justification to be a virulence factor, it is hoped that transgenics carrying the Ubi7/cfp construction will possess a degree of immunity from the Cercospora beticola-induced infection and destruction of mature sugar beet leaves.

Three species of fluorescent *Pseudomonas* were isolated from cultures enriched from the rhizosphere of healthy plants have been bacteriologically cloned and biochemically characterized. One was determined by fatty acid (FA) analysis, done in collaboration with Jeffrey Buyer of the Soil Microbial Systems Lab, to be an isolate of *Pseudomonas syringae*. Two other species, one named *P. corrugata* and another an unidentified *Pseudomonas*, related to *P. putida*, *P. chlororaphis*, *P. corrugata*, *P.migule*, and *P. veronii*. On the basis of FA analysis alone, it appears to be entirely new to bacteriology. The *P.corrugata* and the new *Pseudomonas* species were clearly demonstrated for the first time to produce antibiotics against *Cercospora beticola*.



Figure 1. *Cercospora* on left is strongly inhibited by diffusable substances or antibiotics produced by the new *Pseudomonas* on right.

Pseudomonas spp. strain ND9L will be subjected to 16S RNA sequence analysis to determine phylogenetic relationships of the new species. This is important since these bacteria could be valuable new sources of Cercospora-killing genes for sugar beet bioengineering.

In summary, in this first year of SBDF project #831, several new sugar beet bioengineering stratagies for controlling *Cercospora* were pursued: (1) the pathogen-derived gene *cfp*, conferring resistance to the probable virulence factor cercosporin toxin, has been modified *in vitro* to enhance *in vivo* expression in transgenic sugar beets and confer resistance, (2) together with Joe Saunders, a new method of directly selecting cercosporin resistant cell lines *in vitro* was devised, and (3) beneficial, plant-associated *Pseudomonas* species that exhibit potent antagonism of virulent pathogenic strains of *Cercospora beticola* were discovered and at least one, evidently a new species, could prove to be useful for effectively controlling leafspot when sprayed as a biofungicide either before or during periods of predicted *Cercospora* outbreak. Collaboration with Dr. John Weiland of Fargo, ND is important to implement this latter approach.

Although the bacterial genes involved have yet to be identified or isolated, the construction of sugar beet transgenics that carry the anti-Cercospora genes from Pseudomonas could be a promising approach if sufficient time and funding permit.

We thank Dr. Garry Smith and John Eide for mining the rhizosphere bacteria.

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Characterization of a Fungal Pathogen of the Sugarbeet Root Maggot BSDF Project 850

Chris A. Wozniak (Ann C. Smigocki, Michael R. Marshall)

Publications: Hodge, K.T., Humber, R.A. and Wozniak, C.A. *Cordyceps variabilis* and the genus *Syngliocladium*, Mycologia 90:743-753, 1998.

U.S. Patent US05955071, Fungal Species for the Biological Control of the Sugarbeet Root Maggot, C.A. Wozniak, USDA-ARS; issued 9/21/99.

Abstract: Wozniak, C.A., A novel fungus pathogenic to the sugarbeet root maggot, J Sugarbeet Research 36(3):98, 1999.

Progress Report: As part of an ongoing project to develop biological control agents for management of the sugarbeet root maggot, this project seeks to characterize the fungus *Syngliocladium tetanopsis*. Following its discovery in 1994 in the Red River Valley, *S. tetanopsis* has been assayed for pathogenicity toward the sugarbeet root maggot (SBRM) through *in vitro* assays. All isolates collected (*i.e.*, > 30) have proven infective to SBRM when evaluated against third instar larvae and mortality has been as high as 96 % (n=120) with some isolates. This fungus represents a new species of *fungi imperfecti*; the teleomorphic stage is currently unknown.

Current objectives for research on this agent include the refinement of culture conditions to enhance the rate and quantity of spore production, assess the viability of spore preparations through fluorescent cellular probes, and to determine the host range of *S. tetanopsis*.

In order to develop this fungus for commercial use, attention is paid to the economics of scale-up, including the culture medium needed for spore production. Highly infective spore preparations have been produced using a modification of an oatmeal medium (OatM) as used in standard mycology applications. Although spore yield is high, it would be advantageous to speed up the rate of production. With this in mind, several added sources of organic nitrogen were evaluated for their effects on cultural morphology and growth rate. On OatM, colony morphology is largely restricted to small constricted colonies with a smooth edge and few aerial hyphae. Conidiophores are borne directly on surface hyphae in slime droplets and ultimately on aerial synnemata. Spores from both sources are identical in morphology and they give rise to identical colony types when subcultured; both are also infective to SBRM.

In contrast, when yeast extract, tryptone, casein, potato extract, tomato extract, or complex mixtures of peptone and corn meal with various plant-based carbohydrates were used to amend media, spore production was severely inhibited. Additionally, aerial hyphae and synnemata were absent with a drastically different cultural morphology resulting. It is clear that this species is highly pleomorphic and capable of responding to a variety of nutritional components. Based

upon previous work with alteration of atmospheric conditions of culture (e.g., CO₂, N₂, low O₂) and the current observations, this species is capable of a dimorphic growth habit. While not representing the desired effect (i.e., enhanced spore production), the net growth rate of fungal colonies was greater than with OatM.

Interestingly, of the three isolates evaluated in these studies on media components, one showed a complete lack of growth on modified Sabauraud medium, while the other two showed significant growth, but a lack of sporulation. This suggests that considerable diversity exists within the population(s) of *S. tetanopsis* isolated from Minnesota and North Dakota. Studies on the virulence of these strains toward SBRM will also emphasize examining as many strains as possible to select for variance in these characters.

A lack of spore production was also noted when liquid shake cultures of *S. tetanopsis* were initiated in soy and beef protein digests, although again, the rate of biomass production was rapid. A medium typically used for culture of insect cells was also inoculated with conidiospores of *S. tetanopsis* for assessment of spore production. This medium is rich in animal serum proteins and vitamins, as well as buffering salts. Vegetative growth was observed to progress more rapidly than with potato dextrose broth, however, no spores were observed even after extended culture. A medium consisting of homogenized meal worm larvae was also inoculated with conidiospores to assess the impact of complex insect constituents on growth, however, *S. tetanopsis* grew poorly on this substrate.

Measurements of saline spore suspensions from OatM plates indicated that spore viability was low as measured by hydrolysis of fluorescein diacetate (FDA) by membrane-bound esterases. Serial dilutions onto OatM indicated that spore germination was significantly higher than predicted from FDA measurements, however. Use of FDA and a derivative, carboxyfluorescein diacetate, succinimidyl ester, indicates that these previous assessments of fluorescence may be flawed in that the pH and possibly the ionic strength of the suspending medium were minimizing observed fluorescence. Newly generated protocols indicate that spore viability can be measured with these substrates by adjustment of pH (*i.e.*, increasing alkalinity) and reduction of ionic strength. Serial dilutions offer some information to assess viability, although it appears that there may be an inhibitor of germination present in the spores in that dilutions (plate counts) are nonlinear. Further efforts will address this phenomenon to develop a simple and rapid assay for determination of viability.

Work with cryopreserved spore and mycelial preparations determined that viability could be maintained for at least 16 months at -80°C. More relevant, however, is the stability of preparations as would be typical of biopesticidal products (*i.e.*, shelf-life at room temperature or under refrigeration). Cultures dried under ambient conditions have yielded viable spores after 8 months, although quantitation was not possible at the time of assay. Somewhat surprisingly, spore preparations maintained in 0.85 % saline for 5 months at room temperature yielded viable colonies when plated onto OatM. These findings suggest that spore stability over time may not be a limiting factor in development of a commercial formulation. These experiments will be repeated once the details of the fluorescence viability assays are completed.

While not being pathogenic to the coleopteran, lepidopteran or neuropteran species examined so far, it is plausible that other dipteran species may be within the host range of *S. tetanopsis*. Other members of this genus have been found on flies unrelated to the SBRM. Both *Drosophila melanogaster*, the common fruit fly, and *Musca domestica*, the house fly, are currently being examined for susceptibility to this fungus. Preliminary data indicate that these species are not detrimentally affected by treatment with conidia of *S. tetanopsis*, however, variables associated with the culture of these flies *in vitro* complicate this assessment. More work will be needed to determine this unequivocally. Contacts have also been made with other ARS researchers to test other dipteran species that are considered pests with spore preparations for evaluation of host range.

A recent press release on some of this work by the USDA/ARS Information Staff has garnered some commercial interest in this fungus for root maggot control. Arrangements are being made to share cultures with interested parties and further the evaluation of this potential biopesticide. It is proposed that with the appropriate formulation technology, this agent could be evaluated in the 2001 growing season under field conditions. Most likely this agent would be applied as a granular in-furrow at planting or as a seed coating treatment. Collaborative efforts have been established with ARS and University researchers to evaluate this agent under field conditions of high maggot infestations.

SUGAR BEET RESEARCH 1999 REPORT

Section H

University of Illinois Urbana, Illinois

Dr. D. R. Bush

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by D. R.	. Bush	НЗ

BEET SUGAR DEVELOPMENT FOUNDATION Research Report 2000

New Strategies for Modifying Sucrose Distribution in Sugarbeet

Daniel R. Bush
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The proton-coupled sucrose transport protein mediates the key step in the long-distance transport of newly synthesized sucrose from the leaf to the taproot for storage. While determining how this protein works, we discovered a modified version of the transporter that is 15-fold more active than the wild-type (Lu and Bush 1998). The modified version of the transport protein is an excellent candidate for genetic engineering because it is capable of loading plant cells with molar concentrations of sucrose. Thus, directed expression of the "hyperactive" transporter in the taproot could be used to enhance sucrose accumulation by increasing the uptake capacity of the storage cells. One goal of this project is to test the hypothesis that directed expression of the sucrose transporter can modify sucrose accumulation.

The second aim of this project is to investigate further our recent discovery of a control pathway that regulates sucrose loading into the vascular system in the leaf (Chiou and Bush 1998). The vascular system mediates the long distance transport of sucrose from the photosynthetic cells in the leaf to the sucrose storage cells in the tap root. This was a very significant finding because loading the vascular system for sugar export from the leaf is the key step that determines how much sucrose is delivered to the tap root. Defining the biochemical steps involved in controlling sucrose distribution to the beet will allow us to develop new strategies for manipulating productivity (Bush 1999).

The goal of this project is to increase sucrose storage in the taproot using two approaches. In the first, we are developing transgenic methods to express the hyperactive sucrose transporter in new cells and tissues. The second approach is based on the hypothesis that determining the mechanism the plant uses to control sugar export from the leaf will allow us to develop biochemical and/or biotechnological strategies to increase sucrose transport to the tap root.

Recent Progress

We have been working on the hyperactive form the sucrose transporter that we want to express in the tap root as a mechanism to increase sucrose loading. We have vectors and promoters that should drive expression in this organ. We are now developing collaborations with labs that routinely transform sugar beet to make transgenic plants. Although we generate transgenic Arabidopsis and tobacco using Agrobacterium, the methods for sugar beet are beyond the capability of my lab (beet requires cell culture and we do not have a "gene gun" to deliver the genes). Because beet transformation is moving forward slowly, we are using potato, which is an easily transformed plant with a large storage organ, to test the hypothesis that we can alter sugar accumulation by over-expressing the hyperactive transporter in a target tissue. This is a parallel

experiment that allows us to develop our materials and methods (vectors, growth analysis, measurement of sugars and photosynthesis) while waiting for transformed beets.

The objective of our investigation of the regulatory system that controls sugar export from the leaf is to identify the biochemical steps involved in modifying the sucrose transporters ability to load the vascular tissue of the plant. Our initial analysis of this system showed that it controls sugar allocation between photosynthetic tissues and "import-dependent" organs like the beet tap root (Chiou and Bush 1998). Using Western blot analysis, we recently showed that down regulation of sugar transport activity is the result of protein degradation where the transporter is removed from cells that load the leaf vascular system. In parallel with its turnover, we used nuclear run-offs to show that decrease transporter-mRNA abundance is the result of down-regulation of gene expression. These changes in transporter protein stability and synthesis (as reflected by mRNA abundance) occur within a few hours. Thus, it appears that dynamic regulation of sucrose transporter abundance in the vascular system controls sugar allocation. We are now testing different pharmacological agents as tools for identifying the signal-transduction pathways that participate in this complex control system

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